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A LIMITED LIABILITY PARTNER

10853 U.S. PT 09/692412

Box Patent Application

October 20, 2000

Commissioner for Patents Washington, D.C. 20231

Re: U.S. Non-Provisional Utility Patent Application

Application No.: To Be Assigned

Filed:

Herewith

For:

Plant Polymorphic Markers and Uses Thereof

Inventors:

David F. BUSH et al.

Atty. Docket: 04983.0206.CPUS01/38-21(15493)C

Sir:

The following documents are forwarded herewith for appropriate action by the U.S. Patent and Trademark Office:

- 1. Utility Patent Application Transmittal (PTO/SB/05);
- 2. U.S. Utility Patent Application entitled:

Plant Polymorphic Markers and Uses Thereof

and naming as inventors:

David F. BUSH, Steven D. ROUNSLEY, and Roger C. WIEGAND

the application consisting of:

- a. A specification containing:
  - (i) <u>181</u> pages of a description prior to the claims;
  - (ii) 6 page of claims (26 claims); and
  - (iii) a one (1) page abstract;
- 3. Three (3) CD-ROMs containing the sequence listing; and
- 4. Two (2) return postcards.

This application is being filed without an executed Declaration, and without payment of official fees.



Submitted herewith are a sequence listing on CD-ROM (two copies) and a sequence listing Computer Readable Form (CRF) on CD-ROM, in compliance with 37 C.F.R. §§ 1.52(e), 1.77, AND 1.823 (AS AMENDED IN 65 Fed. Reg. 54603). It is Applicants' understanding that the U.S. Patent and Trademark Office is currently accepting sequence listing submissions in the format prescribed by the amended Code of Federal Regulations, although the new rules do not become mandatory until November 7, 2000.

Three CD-ROMs accompany this transmittal letter: two copies of a sequence listing (Copy 1 and Copy 2) and a sequence listing CRF. Copy 1 and Copy 2 of the sequence listing are identical to each other and to the sequence listing CRF. All three CD-ROMs each contain one file called "Marker Report.rpt" which is 13,491 kilobytes in size and was created on October 19, 2000. These CD-ROMs are IBM-PC machine format, and are MS-DOS and MS-Windows compatible.

It is respectfully requested that, of the two attached postcards, one be stamped with the filing date of these documents and returned to our courier, and the other, prepaid postcard, be stamped with the filing date and unofficial application number and returned as soon as possible.

Respectfully submitted,

David R. Marsh (Reg. No. 41,408) June E. Cohan (Reg. No. 43,741)

Enclosures

PTO/SB/05 (2/98) Approved for use through 09/30/00. OMB 0651-0032

Patent and Trademark Office: U.S. DEPARTMENT OF COMMERCE
Under the Paperwork Reduction Act of 1995, no persons are required to respond to a collection of information unless it displays a valid OMB control number.

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Signature					Date	October 20,	October 20, 2000		

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# PLANT POLYMORPHIC MARKERS AND USES THEREOF FIELD OF THE INVENTION

The present invention is in the field of plant genetics. More specifically, the invention relates to nucleic acid markers associated with *Arabidopsis thaliana* ecotypes.

5 The invention also relates to methods for detecting polymorphisms.

# INCORPORATION OF SEQUENCE LISTING

This application contains a sequence listing, which is contained on three identical CD-ROMs: two copies of a sequence listing (Copy 1 and Copy 2) and a sequence listing Computer Readable Form (CRF), all of which are herein incorporated by reference. All three CD-ROMs each contain one file called "Marker Report.rpt" which is 13,491 kilobytes in size and was created on October 19, 2000.

#### **BACKGROUND OF THE INVENTION**

## I. Arabidopsis thaliana

The identification in *Arabidopsis thaliana* of polymorphic markers is important in the development of nutritionally enhanced or agriculturally enhanced crops. Such polymorphic markers are useful in, for example, genetic mapping or linkage analysis, marker assisted breeding, physical genome mapping, transgenic crop production, crop monitoring diagnostics, and gene identification and isolation.

Arabidopsis thaliana is widely used as a model organism for basic and applied research in the biology of flowering plants. Arabidopsis thaliana is a model system for plant genomic research in part due to its small and characterized genome, which is estimated to be comprised of approximately 20,000 to 25,000 genes. The genome is estimated to have a haploid content of around 100Mb, present on five chromosomes. Reported partial sequence analysis has provided information on genome features such as gene density and gene structure (Settles and Byrne, Genome Research 8:83-85 (1998), the

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entirety of which is herein incorporated by reference). Based on reports from the European Union Sequencing Consortium, the average gene density is one gene every approximately 4.8kb.

Other important characteristics that make Arabidopsis thaliana a useful test system include its rapid life-cycle, small size, which allows for controlled growth in restricted space, its prolific seed production, the availability of characterized and uncharacterized mutants and the existence of a reliable transformation system.

Molecular genetics is often used in the analysis of plant genes and is particularly useful in the analysis of complex biological processes such as developmental regulation. In one approach the use of mutant plants, e.g. Arabidopsis thaliana mutants, in molecular genetic research requires the location of the mutation. Molecular markers are a useful way to locate such mutations.

Identification of target loci and the isolation of associated genes using molecular markers has been reported (Liu et al., Proc. Natl. Acad. Sci. USA, 96:6535-6540 (1999); Muramoto et al., The Plant Cell, 11:335-347 (1999); Bowman and Smyth, Development, 126:2387-2396 (1999); Michaels and Amasino, The Plant Cell, 11:949-956 (1999); Ha et al., The Plant Cell, 11:1153-1163 (1999); Walker et al., The Plant Cell, 11:1337-1349 (1999); Sedbrook et al., Proc. Natl. Acad. Sci. USA, 96:1140-1145 (1999); Kiyosue et al., Proc. Natl. Acad. Sci. USA, 96:4186-4191 (1999); and Davis et al., Proc. Natl. Acad. Sci. USA, 96:6541-6546 (1999), all of which are herein incorporated by reference in their entirety). The use of markers to isolate a genomic region of interest is often referred to as map based cloning, chromosome walking or positional cloning. Many of the Arabidopsis thaliana markers that have been used in map based cloning are anchored to genetic maps such as the Lister & Dean map (See e.g. http://genome-

www3.stanford.edu/cgi-bin/AtDB/RIintromap).

Physical or partial physical maps of the Arabidopsis thaliana genome have also been reported (See e.g. http://genome-www3.stanford.edu/atdb\_welcome.html). A

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physical map of *Arabidopsis thaliana*, Columbia based on a collection of bacterial artificial chromosomes (BACs) is available (Marra *et al.*, *Nat. Genet.*, 22(3):265-270 (1999); Mozo *et al.*, *Nat. Genet.*, 22(*e*):271-275 (1999), both of which are herein incorporated by reference in their entirety). An overlapping series of BACs representing the *Arabidopsis thaliana*, Columbia genome is available from AIMS, Arabidopsis Biological Resource Center, 309 B&Z Building, 1735 Neil Avenue, Columbus, OH 43210, USA.

Cho et al. reported a low density biallelic polymorphic map based on a comparison of Arabidopsis thaliana, Columbia and Arabidopsis thaliana, Landsberg erecta ecotypes by screening approximately 0.5% of the genome for such polymorphisms (Cho et al., Nature Genetics 23:203-207 (1999), the entirety of which is herein incorporated by reference). In this survey 487 single nucleotide polymorphisms (SNPs) were reported. Cho et al. also reported the use of oligonucleotide arrays to detect Arabidopsis thaliana SNPs.

The present invention provides polymorphic nucleic acid markers whose physical location is known within the *Arabidopsis thaliana* genome. Moreover, the physical location of such markers is further known within a particular BAC and the position of that BAC relative to other BACs in the genome is also known.

Successful isolation of a region of *Arabidopsis thaliana* DNA associated with a trait of interest requires a nucleic acid marker to be sufficiently close to the trait. As the present invention provides a collection of nucleic acid markers in the *Arabidopsis thaliana* genome which allows for the efficient isolation of regions of *Arabidopsis thaliana* DNA associated with traits of interest. Moreover, the association of a collection of nucleic acid markers with a trait of interest may be simultaneously investigated.

### **Summary of the Invention**

The present invention provides a collection of nucleic acid molecules capable of detecting a set of polymorphisms as shown in Table A.

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The present invention also includes and provides a method of isolating a region of genomic DNA associated with a phenotype of interest comprising: (A) identifying an *Arabidopsis* plant having a first ecotype with the phenotype; (B) crossing the *Arabidopsis* plant with an *Arabidopsis* plant having a second ecotype lacking the phenotype of interest; (C) propagating and self pollinating seeds from the cross; (D) selecting progeny of self pollinated seeds with the phenotype of interest; (E) screening progeny of self pollinated seeds with the phenotype of interest with a collection of nucleic acid molecules, the collection of nucleic acid molecules capable of detecting a set of polymorphisms where the polymorphisms are distributed throughout the genome of the self pollinated seeds with the phenotype of interest at an average density of more than one polymorphism per about 100kb; (F) calculating the linkage of each of the nucleic acid molecules to the phenotype; and (G) isolating said region of genomic DNA associated with the phenotype based on its linkage to one or more of the nucleic acid molecules.

The present invention also provides a method of identifying a region of genomic DNA associated with a phenotypic trait of interest comprising: (A) screening a mapping population of *Arabidopsis* plants to determine the linkage of the phenotypic trait with a collection of nucleic acid molecules, wherein the nucleic acid molecules are capable of detecting a set of polymorphisms, where the polymorphisms are distributed throughout the genome of the mapping population of *Arabidopsis* plants at an average density of more than one polymorphism per about 100kb; (B) calculating the linkage of each of the nucleic acid molecules to the phenotypic trait; and (C) identifying the genomic DNA region associated the phenotypic trait based on its linkage to one or more of the nucleic acid molecules.

The present invention also provides a method of identifying a nucleic acid molecule associated with a phenotypic trait comprising: (A) screening a mapping population of *Arabidopsis* plants to determine the linkage of the phenotypic trait with polymorphisms, wherein the polymorphisms are distributed throughout the genome of the

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mapping population of *Arabidopsis* plants at an average density of more than one polymorphism per about 100kb; (B) calculating the linkage of each of the polymorphism to the phenotypic trait; and (C) isolating the nucleic acid molecule associated with the phenotypic trait based on its linkage to one or more of the polymorphisms.

The present invention also provides a method of isolating a nucleic acid molecule associated with a phenotypic trait comprising: (A) screening a mapping population of *Arabidopsis* plants to determine the linkage of the phenotypic trait with a polymorphism, wherein the polymorphism is selected from the group consisting of a polymorphism from Table A; and (B) isolating the nucleic acid molecule associated with the phenotypic trait based on its linkage to one or more of the polymorphisms.

The present invention also provide a method of introgressing a trait of interest into a plant comprising using a nucleic acid marker for marker assisted selection of the plant, the nucleic acid marker capable of detecting a polymorphism selected from Table A, and introgressing the trait into said plant.

The present invention also provides a collection of non-identical nucleic acid molecules capable of detecting polymorphisms present in an *Arabidopsis* mapping population, wherein the collection of non-identical nucleic acid molecules is capable of detecting at least 25 polymorphisms selected from the group consisting of Table A.

The present invention also provides a computer readable medium having recorded thereon at least 100 of the polymorphisms set forth in Table A.

The present invention also provides a method for identifying transposons in the DNA of an organism comprising identifying INDELs in that DNA and comparing the sequence of the INDELs to the sequence of one or more known transposons.

#### **Detailed Description of the Invention**

The genomes of animals and plants naturally undergo spontaneous mutation in the course of their continuing evolution (Gusella, *Ann. Rev. Biochem. 55*:831-854 (1986), the entirety of which is herein incorporated by reference). A "polymorphism" is a variation

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or difference in the sequence of a genetic region that arises in some of the members of a species. Variant sequences can be defined with reference to an arbitrary or non-arbitrary standard sequence for the species. A polymorphism is thus said to be "allelic," in that, due to the existence of the polymorphism, some members of a species may have the "standard" sequence (*i.e.* the standard "allele") whereas other members may have a variant sequence (*i.e.*, a variant "allele"). Thus, as used herein, an allele is one of two or more alternative versions of a gene or other genetic region at a particular location on a chromosome. In the simplest case, only one variant sequence may exist, and the polymorphism is thus said to be bi-allelic. In other cases, the species' population may contain multiple alleles, and the polymorphism is termed tri-allelic, *etc*.

A single gene or genetic region may have multiple different unrelated polymorphisms. For example, it may have a one bi-allelic polymorphism at one site, another bi-allelic polymorphism at another site and a multi-allelic polymorphism at another site. When all the sequences for a group of alleles at a chromosomal locus in a plant are the same, the alleles are said to be "homozygous" at that locus. When the sequence of any allele at a particular locus in a plant is different, the population of alleles is said to be "heterozygous" at that locus.

Phenotypic traits can vary due to environmental and/or genetic factors. For example, polymorphisms at a particular chromosomal locus can affect the phenotypic trait associated with that locus.

As used herein, a phenotypic trait of interest may be any trait exhibited by a plant, whether naturally occurring or otherwise, that is capable of being inherited. Moreover, the phenotypic trait of interest may, for example, be transient, permanent or only present when the plant or part thereof is subjected to environmental stimuli or challenge. A phenotypic trait of interest may be a desired trait. In other cases the phenotypic trait of interest may be an undesired trait. Furthermore, phenotypic traits are not limited to visible traits. While the phenotypic trait may be any trait, preferred traits of interest are those

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that have agricultural significance. Examples of agricultural traits include those that affect a component of yield, those that provide disease or chemical resistance, and those that affect developmental traits such as pollen or ovule production, *etc.*, and those that affect composition of plants or plant parts, including seed proteins or oils, starch or sugar composition, nutrient content and the like.

Many phenotypic traits are the result of multiple genes or genetic factors, for example, a phenotypic trait that is the result of a quantitative trait allele. An allele of a quantitative trait locus (QTL) can, of course, comprise multiple genes or other genetic factors even within a contiguous genomic region or linkage group. As used herein, an allele of a quantitative trait locus can therefore encompass more than one gene or other genetic factor where each individual gene or genetic component is also capable of exhibiting allelic variation and where each gene or genetic factor also has a phenotypic affect on the quantitative trait in question.

As used herein, a "marker" is an indicator for the presence of at least one polymorphism. A marker is preferably a nucleic acid molecule. It is understood that a marker can, for example, be an oligonucleotide probe or primer.

A "nucleic acid marker" as used herein means a nucleic acid molecule that is capable of being a marker for detecting a polymorphism.

useful, *e.g.* for hybridizing probes, nucleotide array elements or amplification primers.

Oligonucleotide molecules are comprised of two or more nucleotides, *i.e.*deoxyribonucleotides or ribonucleotides, preferably more than five and up to 30 or more.

The exact size will depend on many factors, which in turn depend on the ultimate function or use of the oligonucleotide. Oligonucleotides can comprise ligated natural

nucleic molecules acids or synthesized nucleic acid molecules and comprise between 5 to 150 nucleotides or between about 15 and about 100 nucleotides, or preferably up to 100 nucleotides, and even more preferably between 15 to 30 nucleotides or most preferably

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between 18-25 nucleotides, identical or complementary to a sequence of similar length. This invention provides oligonucleotides capable of detecting polymorphisms. Such oligonucleotides may be nucleic acid elements for use on solid arrays (e.g. synthesized or spotted). Such oligonucleotides may also be primers for use in polymerase chain reaction (PCR) or other reactions. The term "primer" as used herein refers to a nucleic acid molecule, preferably an oligonucleotide whether derived from a naturally occurring molecule such as one isolated from a restriction digest or one produced synthetically, which is capable of acting as a point of initiation of synthesis when placed under conditions in which synthesis of a primer extension product which is complementary to a nucleic acid strand is induced, i.e., in the presence of nucleotides and an agent for polymerization such as DNA polymerase and at a suitable temperature and pH. The primer is preferably single stranded for maximum efficiency in amplification, but may alternatively be double stranded. If double stranded, the primer is first treated to separate its strands before being used to prepare extension products. Preferably, the primer is an oligodeoxyribonucleotide. The primer must be sufficiently long to prime the synthesis of extension products in the presence of the agent for polymerization. The exact lengths of the primers will depend on many factors, including temperature and source of primer. For example, depending on the complexity of the target sequence, the oligonucleotide primer typically contains at least 15, more preferably 18 nucleotides, which are identical or complementary to the template and optionally a tail of variable length which need not match the template. The length of the tail should not be so long that it interferes with the recognition of the template. Short primer molecules generally require cooler temperatures to form sufficiently stable hybrid complexes with the template.

The primers herein are selected to be "substantially" complementary to the different strands of each specific sequence to be amplified. This means that the primers must be sufficiently complementary to hybridize with their respective strands. Therefore, the primer sequence need not reflect the exact sequence of the template. For example, a

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non-complementary nucleotide fragment may be attached to the 5' end of the primer, with the remainder of the primer sequence being complementary to the strand. Alternatively, non-complementary bases or longer sequences can be interspersed into the primer, provided that the primer sequence has sufficient complementarity with the sequence of the strand to be amplified to hybridize therewith and thereby form a template for synthesis of the extension product of the other primer. Computer generated searches using programs such as Primer3 (<a href="www-genome.wi.mit.edu/cgi-bin/primer/primer3.cgi">www-genome.wi.mit.edu/cgi-bin/primer/primer3.cgi</a>), STSPipeline (<a href="www-genome.wi.mit.edu/cgi-bin/www-STS\_Pipeline">www-genome.wi.mit.edu/cgi-bin/www-STS\_Pipeline</a>), or GeneUp (Pesole et al., BioTechniques 25:112-123 (1998), the entirety of which is herein incorporated by reference), for example, can be used to identify potential PCR primers. Exemplary primers include primers that are 18 to 50 bases long, where at least between 18 to 25 bases are identical or complementary to at least 18 to 25 bases of a segment of the template sequence.

This invention also contemplates and provides primer pairs for amplification of nucleic acid molecules in order to detect polymorphisms. As used herein "primer pair" means a set of two oligonucleotide primers based on two separated sequence segments of a target nucleic acid sequence. One primer of the pair is a "forward primer" or "5' primer" having a sequence which is identical to the more 5' of the separated sequence segments (+ strand). The other primer of the pair is a "reverse primer" or "3' primer" having a sequence which is complementary to the more 3' of the separated sequence segments (+ strand). A primer pair allows for amplification of the nucleic acid sequence between and including the separated sequence segments. Optionally, each primer pair can comprise additional sequences, *e.g.* universal primer sequences or restriction endonuclease sites, at the 5' end of each primer, *e.g.* to facilitate cloning, DNA sequencing, or reamplification of the target nucleic acid sequence.

As used herein, a "mapping population" is a collection of plants capable of being used with markers to map the genetic position of traits.

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As used herein, a polymorphic marker is a marker capable of detecting one or more polymorphisms.

The present invention provides nucleic acid molecules which are markers, *i.e.* capable of detecting polymorphisms that are distributed throughout the genome of a mapping population.

As used herein, a "characterized polymorphism" is a polymorphism whose physical position on a genome is known. In a preferred embodiment, the physical position of a characterized polymorphism on an isolated nucleic acid molecule, such as a bacterial artificial chromosome comprising *Arabidopsis thaliana* genomic DNA, is known. Thus the present invention also provides nucleic acid molecules capable of detecting characterized polymorphisms throughout a genome.

In a further preferred embodiment, a characterized polymorphism is any polymorphism where the nucleic acid sequences of at least two of the polymorphisms present in an *Arabidopsis* mapping population are known (sequenced characterized polymorphism). In a particularly preferred embodiment, a characterized polymorphism from Table A. In another particularly preferred embodiment, a characterized polymorphism from Table A is part of a collection of polymorphisms, where preferably over 25%, more preferably over 50% and even more preferably over 75% of the polymorphisms are selected from the polymorphisms in Table A.

The present invention provides nucleic acid molecules capable of detecting insertion/deletion polymorphisms (INDELs) in *Arabidopsis* at an average density of one INDEL per 8.4 kb. The present invention also provides nucleic acid molecules capable of detecting single nucleotide polymorphisms (SNPs) at an average density of one SNP per 3.9 kb. The present invention also provides nucleic acid molecules capable of detecting polymorphisms at an average density of one polymorphism per 2.7 kb.

As used herein, an "INDEL" is any insertion/deletion polymorphism characterized by additional nucleotides in at least one allele as compared to a reference allele. As used

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herein, a "SNP" is any polymorphism characterized by a different single nucleotide at a particular physical position in at least one allele.

The polymorphisms capable of detection by nucleic acid molecules of the present invention are distributed throughout the genome of the mapping population in a manner that allows the efficient identification of a genomic region associated with a phenotypic trait. In a preferred embodiment, the polymorphisms are distributed throughout the genome where 60%, preferably 70%, more preferably 80%, even more preferably 90%, 95% or 100% of the genome has a characterized polymorphism at a density of higher than one polymorphism per 100kb, more preferably higher than one polymorphism per 50kb, and even more preferably higher than one polymorphisms are distributed throughout the genome where 60%, preferably 70%, more preferably 80%, even more preferably 90%, 95% or 100% of genome has a characterized polymorphism at a density of higher than one polymorphism per 3.5cM, more preferably higher than one polymorphism per 3.5cM, and even more preferably higher than one polymorphism per 3.25cM, and even more preferably higher than one polymorphism per 3.0cM, 2.75cM, 2.5cM, 2.0cM, 1.5cM, 1.0cM or 0.5cM.

In a preferred embodiment of the present invention, the efficient identification of a genomic region associated with a phenotypic trait, *e.g.* a QTL or a single gene, is provided, where the genomic region is less than 100kb, more preferably less than 50kb, and even more preferably less than 25kb, 10kb, 7kb, 5kb or 3kb from a characterized polymorphism. In another preferred embodiment of the present invention the efficient identification of a genomic region associated with a phenotypic trait where the genomic region is less than 3.5cM, more preferably less than 3.25cM, and even more preferably less than 3.75cM, 2.75cM, 2.5cM, 2.0cM, 1.5cM, 1.0cM or 0.5cM from a characterized polymorphism.

It is understood that the distribution of polymorphisms need not be uniform in a genome as certain regions will exhibit a higher average density of polymorphisms (e.g.

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non-centromeric regions) and certain regions will exhibit a lower average density of polymorphisms (e.g. centromeric regions).

In a preferred embodiment, the efficient identification of a genomic region associated with a phenotypic trait of interest will be obtained by a simultaneous screening for the presence of 25 or more, more preferably 50 or more, even more preferably 75 or more, 100 or more, 150 or more, 200 or more, 250 or more, 300 or more, 400 or more or 500 or more, 1,000 or more, 2,000 or more, 3,000 or more, 4,000 or more polymorphisms. In an even more preferred embodiment, the efficient identification of a genomic region associated with a phenotypic trait of interest will be obtained by a simultaneously screening for the presence of 25 or more, more preferably 50 or more, even more preferably (where appropriate) 100 or more, or 250 or more *etc.* of the polymorphisms in Table A.

In another preferred embodiment, the efficient identification of a genomic region associated with a phenotypic trait of interest will be obtained by screening for the presence of 25 or more, more preferably 50 or more, even more preferably 75 or more, 100 or more, 150 or more, 200 or more, 250 or more, 300 or more, 400 or more or 500 or more, 1,000 or more, 2,000 or more, 3,000 or more, 4,000 or more polymorphisms during a single assay. In an even more preferred embodiment the efficient identification of a genomic region associated with a phenotypic trait of interest will be obtained by screening for the presence of 25 or more, more preferably 50 or more, even more preferably (where appropriate) 100 or more or 250 or more *etc.* of the polymorphisms in Table A during a single assay. A single assay can comprise many steps. One or more of these steps can occur sequentially.

In an embodiment of the present invention, the assay is carried out using a high throughput system. A particularly preferred high throughput system involves a solid phase array. A particularly preferred solid phase array is a microarray.

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In the assays below, a collection of markers for polymorphisms can comprise from a few up to millions of different nucleic acid molecules. For example, using simple dot-blot hybridization methods, membranes with many nucleic acid molecules can be generated for screening. The solid-phase techniques described below and known in the art can be adapted for high-throughput monitoring of polymorphisms. In such methods different immobilized nucleic acid molecule probes can be placed on a solid support at microarray densities of up to millions of nucleic acid molecules per square inch. Similarly, very large sets of nucleic acid molecules can be immobilized for simultaneous screening against one or more probes.

Several methods have been described for fabricating microarrays of nucleic acid molecules and using such microarrays in detecting nucleic acid sequences. For instance, microarrays of markers for polymorphisms can be fabricated by spotting nucleic acid molecules, e.g. oligonucleotides, onto substrates or fabricating oligonucleotide sequences in situ on a substrate. Spotted or fabricated nucleic acid molecules can be applied in a high density matrix pattern of up to about 30 non-identical nucleic acid molecules per square centimeter or higher, e.g. up to about 100 or even 1,000 per square centimeter or higher. Useful substrates for arrays include nylon, glass and silicon. See, for instance,  $5,202,231;\ 5,242,974;\ 5,384,261;\ 5,405,783;\ 5,412,087;\ 5,424,186;\ 5,429,807;\ 5,436,327;$  $5,445,934;\,5,472,672;\,5,525,464;\,5,527,681;\,5,529,756;\,5,532,128;\,5,545,531;\,5,554,501;$  $5,556,752;\,5,561,071;\,5,571,639;\,5,593,839;\,5,599,695;\,5,624,711;\,5,658,734;\,5,700,637;$ 5,744,305; 5,800,992; 6,004,755 and 6,087,102 the disclosures of all of which are incorporated herein by reference in their entireties. Sequences can be efficiently analyzed by hybridization or primer extension. See, for instance, U.S. Patents 5,202,231;  $5,445,934;\,5,492,806;\,5,525,464;\,5,695,940;\,5,700,637;\,5,744,305;\,5,800,992;\,5,807,522;$ and 5,830,645, all of which are incorporated herein by reference in their entirety. Nucleic acid molecule microarrays may be screened with molecules or fragments thereof to determine nucleic acid molecules that specifically bind molecules or fragments thereof.

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In a preferred embodiment, a microarray of the present invention comprises at least 10 nucleic acid molecules that specifically hybridize under high stringency to at least 10 polymorphic nucleic acid sequences characterized by this invention. In a more preferred embodiment, a microarray of the present invention comprises at least 100 nucleic acid molecules that specifically hybridize under high stringency to at least 100 characterized polymorphic nucleic acid sequences; more preferably at least 1,000 or 2,500 marker nucleic acid molecules that specifically hybridize under high stringency to at least 1,000 or 2,500 characterized polymorphic nucleic acid sequences; even more preferably at least at least 4,000 or more marker nucleic acid molecules that specifically hybridize under high stringency to at least 4,000 or more characterized polymorphic nucleic acid sequences.

In a preferred embodiment, a microarray of the present invention comprises at least 10 nucleic acid molecules capable of detecting or characterizing by primer extension to at least 10 polymorphic nucleic acid sequences characterized by this invention. In a more preferred embodiment, a microarray of the present invention comprises at least 100 nucleic acid molecules capable of detecting or characterizing by primer extension to at least 100 characterized polymorphic nucleic acid sequences; even more preferably at least 1,000 or 2,500 nucleic acid molecules capable of detecting or characterizing by primer extension to at least 1,000 or 2,500 characterized polymorphic nucleic acid sequences; even more preferably at least 4,000 or more nucleic acid molecules capable of detecting or characterizing by primer extension to at least 4,000 or more nucleic acid molecules capable of detecting or characterizing by primer extension to at least 4,000 or more characterized polymorphic nucleic acid sequences.

In a preferred embodiment, the microarray is a variant detector array (VDA)(Cho et al., Nature Genetics 23:203-207 (1999); Wang et al., Science 280: 1077-1082 (1998), the entirety of which is herein incorporated by reference; Winzeler et al., Curr. Opin. Genet. Dev. 4: 602-608 (1997), the entirety of which is herein incorporated by reference). For example, each detection block can consist of four variant detector arrays (VDAs)

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corresponding to the alternative alleles: two for the forward strand sequence and two for the reverse strand sequence (*See e.g.* Cho *et al.*, *Nature Genetics 23:*203-207 (1999)). For each of the interrogated positions (for example, -5 to +5 relative to the polymorphic position), a set of four suitable length oligonucleotides per SNP or other polymorphism (*e.g.* 25-mers are prepared where the oligonucleotides are complementary to the SNP or other polymorphic region except at the interrogated position). Hybridization of the oligonucleotides with the matching allele results in a strong signal.

The detection or screening of polymorphic nucleic acid sites in a sample of DNA may be facilitated, for example, through including the use of nucleic acid amplification methods. Such methods specifically increase the concentration of polynucleotides that span the polymorphic site, or include that site and sequences located either distal or proximal to it. Such amplified molecules can be readily detected by gel electrophoresis or other means.

If a polymorphism creates or destroys a restriction endonuclease cleavage site, or if it results in the loss or insertion of DNA (e.g., a Variable Number of Tandem Repeats (VNTR) polymorphism), it will alter the size or profile of the DNA fragments that are generated by digestion with that restriction endonuclease. As such, individuals that possess a variant sequence can be distinguished from those having the original sequence by restriction fragment analysis. Polymorphisms that can be identified in this manner are termed "restriction fragment length polymorphisms" ("RFLPs"). RFLPs have been widely used in human and plant genetic analyses (Glassberg, UK Patent Application 2135774; Skolnick et al., Cytogen. Cell Genet. 32:58-67 (1982); Botstein et al., Ann. J. Hum. Genet. 32:314-331 (1980); Fischer et al., PCT Application WO 90/13668; Uhlen, PCT Application WO 90/1369, all of which are herein incorporated by reference in their entirety).

An alternative method of determining polymorphisms is based on cleaved amplified polymorphic sequences (CAPS) (Konieczny, A. and F.M. Ausubel, *Plant J.* 

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4:403-410 (1993); Lyamichev et al., Science 260:778-783 (1993), the entireties of which are herein incorporated by reference). One advantage of this method is the large amount of target DNA that is generated by amplification which eliminates the requirement for radiolabeling for detection of the polymorphism.

Polymorphisms can also be identified by single strand conformation polymorphism (SSCP) analysis. The SSCP technique is a method capable of identifying most sequence variations in a single strand of DNA, typically between 150 and 250 nucleotides in length (Elles, Methods in Molecular Medicine: Molecular Diagnosis of Genetic Diseases, Humana Press (1996), the entirety of which is herein incorporated by reference; Orita et al., Genomics 5:874-879 (1989), the entirety of which is herein incorporated by reference). Under denaturing conditions a single strand of DNA will adopt a conformation that is uniquely dependent on its sequence. This conformation usually will be different even if only a single base is changed. Most conformations have been reported to alter the physical configuration or size sufficiently to be detectable by electrophoresis. A number of protocols have been described for SSCP including, but not limited to Lee et al., Anal. Biochem. 205:289-293 (1992), the entirety of which is herein incorporated by reference; Suzuki et al., Anal. Biochem. 192:82-84 (1991), the entirety of which is herein incorporated by reference; Lo et al., Nucleic Acids Research 20:1005-1009 (1992), the entirety of which is herein incorporated by reference; Sarkar et al., Genomics 13:441-443 (1992), the entirety of which is herein incorporated by reference).

Polymorphisms may also be detected using a DNA fingerprinting technique called amplified fragment length polymorphism (AFLP), which is based on the selective PCR amplification of restriction fragments from a total digest of genomic DNA to profile that DNA. Vos *et al.*, *Nucleic Acids Res.* 23:4407-4414 (1995), the entirety of which is herein incorporated by reference. This method allows for the specific co-amplification of many restriction fragments, which can be analyzed without knowledge of the nucleic acid sequence. AFLP employs basically three steps. Initially, a sample of genomic DNA is

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cut with restriction enzymes and oligonucleotide adapters are ligated to the restriction fragments of the DNA. The restriction fragments are then amplified using PCR by using the adapter and restriction sequence as target sites for primer annealing. The selective amplification is achieved by the use of primers that extend into the restriction fragments, amplifying only those fragments in which the primer extensions match the nucleotide flanking the restriction sites. These amplified fragments are then visualized on a denaturing polyacrylamide gel (Beismann et al., Mol. Ecol. 6:989-993 (1997); Janssen et al., Int. J. Syst. Bacteriol 47:1179-1187 (1997); Huys et al., Int. J. Syst. Bacteriol. 47:1165-1171 (1997); McCouch et al., Plant Mol. Biol. 35:89-99 (1997); Nandi et al., Mol. Gen. Genet. 255:1-8 (1997); Cho et al., Genome 39:373-378 (1996); Simons et al., Genomics 44:61-70 (1997); Cnops et al., Mol. Gen. Genet. 253:32-41 (1996); Thomas et al., Plant J. 8:785-794 (1995), all of which are herein incorporated by reference in their entirety).

Polymorphisms may also be detected using random amplified polymorphic DNA (RAPD) (Williams *et al.*, *Nucl. Acids Res. 18*:6531-6535 (1990), the entirety of which is herein incorporated by reference).

SNPs generally occur at greater frequency than other polymorphic markers and are spaced with a greater uniformity throughout a genome than other reported forms of polymorphism. The greater frequency and uniformity of SNPs means that there is greater probability that such a polymorphism will be found near or in a genetic locus of interest than would be the case for other polymorphisms. SNPs are located in protein-coding regions and noncoding regions of a genome. Some of these SNPs may result in defective or variant protein expression (*e.g.*, as a result of mutations or defective splicing). Analysis (genotyping) of characterized SNPs can require only a plus/minus assay rather than a lengthy measurement, permitting easier automation.

SNPs can be characterized using any of a variety of methods. Such methods include the direct or indirect sequencing of the site, the use of restriction enzymes

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(Botstein et al., Am. J. Hum. Genet. 32:314-331 (1980), the entirety of which is herein incorporated reference; Konieczny and Ausubel, Plant J. 4:403-410 (1993), the entirety of which is herein incorporated by reference), enzymatic and chemical mismatch assays (Myers et al., Nature 313:495-498 (1985), the entirety of which is herein incorporated by reference), allele-specific PCR (Newton et al., Nucl. Acids Res. 17:2503-2516 (1989), the entirety of which is herein incorporated by reference; Wu et al., Proc. Natl. Acad. Sci. USA 86:2757-2760 (1989), the entirety of which is herein incorporated by reference), ligase chain reaction (Barany, Proc. Natl. Acad. Sci. USA 88:189-193 (1991), the entirety of which is herein incorporated by reference), single-strand conformation polymorphism analysis (Labrune et al., Am. J. Hum. Genet. 48: 1115-1120 (1991), the entirety of which is herein incorporated by reference), single base primer extension (Kuppuswamy et al., Proc. Natl. Acad. Sci. USA 88:1143-1147 (1991), Goelet US 6,004,744; Goelet 5,888,819; all of which are herein incorporated by reference in their entirety ), solid-phase ELISA-based oligonucleotide ligation assays (Nikiforov et al., Nucl. Acids Res. 22:4167-4175 (1994), dideoxy fingerprinting (Sarkar et al., Genomics 13:441-443 (1992), the entirety of which is herein incorporated by reference), oligonucleotide fluorescencequenching assays (Livak et al., PCR Methods Appl. 4:357-362 (1995a), the entirety of which is herein incorporated by reference), 5'-nuclease allele-specific hybridization TaqMan<sup>™</sup> assay (Livak et al., Nature Genet. 9:341-342 (1995), the entirety of which is herein incorporated by reference), template-directed dye-terminator incorporation (TDI) assay (Chen and Kwok, Nucl. Acids Res. 25:347-353 (1997), the entirety of which is herein incorporated by reference), allele-specific molecular beacon assay (Tyagi et al., Nature Biotech. 16: 49-53 (1998), the entirety of which is herein incorporated by reference), PinPoint assay (Haff and Smirnov, Genome Res. 7: 378-388 (1997), the entirety of which is herein incorporated by reference), dCAPS analysis (Neff et al., Plant J. 14:387-392 (1998), the entirety of which is herein incorporated by reference), pyrosequencing (Ronaghi et al, Analytical Biochemistry 267:65-71 (1999); Ronaghi et al

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PCT application WO 98/13523; Nyren *et al* PCT application WO 98/28440, all of which are herein incorporated by reference in their entirety; http://www.pyrosequencing.com), using mass spectrometry *e.g.*, the Masscode ™ system (Howbert *et al* WO 99/05319; Howber *et al* WO 97/27331, all of which are herein incorporated by reference in their entirety; http://www.rapigene.com; Becker *et al* PCT application WO 98/26095; Becker *et al* PCT application; WO 98/12355; Becker *et al* PCT application WO 97/33000; Monforte *et al* US 5,965,363, all of which are herein incorporated by reference in their entirety), invasive cleavage of oligonucleotide probes (Lyamichev *et al Nature Biotechnology 17*:292-296, herein incorporated by reference in its entirety; http://www.twt.com), using high density oligonucleotide arrays (Hacia *et al Nature Genetics 22*:164-167; herein incorporated by reference in its entirety; http://www.affymetrix.com).

INDELs are identified by comparing sequence of *Arabidopsis thaliana* ecotypes Columbia and Landsberg erecta. Certain INDELs are believed to have resulted from insertion or excision of transposable elements. Thus, INDEL sequences can be used to identify candidate sequences for active transposons by comparing INDEL sequences to the sequence of known transposons. For instance, certain INDEL sequences of greater than 100 bp were found to exhibit similarity to the sequence of MuDR transposable element from maize.

Polymorphisms may also be detected using allele-specific oligonucleotides (ASO), which, can be for example, used in combination with hybridization based technology including southern, northern, and dot blot hybridizations, reverse dot blot hybridizations and hybridizations performed on microarray and related technology.

The stringency of hybridization for polymorphism detection is highly dependent upon a variety of factors, including length of the allele-specific oligonucleotide, sequence composition, degree of complementarity (*i.e.* presence or absence of base mismatches), concentration of salts and other factors such as formamide, and temperature. These

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factors are important both during the hybridization itself and during subsequent washes performed to remove target polynucleotide that is not specifically hybridized. In practice, the conditions of the final, most stringent wash are most critical. In addition, the amount of target polynucleotide that is able to hybridize to the allele-specific oligonucleotide is also governed by such factors as the concentration of both the ASO and the target polynucleotide, the presence and concentration of factors that act to "tie up" water molecules, so as to effectively concentrate the reagents (*e.g.*, PEG, dextran, dextran sulfate, *etc.*), whether the nucleic acids are immobilized or in solution, and the duration of hybridization and washing steps.

Hybridizations are preferably performed below the melting temperature  $(T_m)$  of the ASO. The closer the hybridization and/or washing step is to the  $T_m$ , the higher the stringency.  $T_m$  for an oligonucleotide may be approximated, for example, according to the following formula:  $T_m = 81.5 + 16.6 \times (\log 10[\text{Na+}]) + 0.41 \times (\% G+\text{C}) - 675/\text{n}$ ; where [Na+] is the molar salt concentration of Na+ or any other suitable cation and n = number of bases in the oligonucleotide. Other formulas for approximating  $T_m$  are available and are known to those of ordinary skill in the art.

Stringency is preferably adjusted so as to allow a given ASO to differentially hybridize to a target polynucleotide of the correct allele and a target polynucleotide of the incorrect allele. Preferably, there will be at least a two-fold differential between the signal produced by the ASO hybridizing to a target polynucleotide of the correct allele and the level of the signal produced by the ASO cross-hybridizing to a target polynucleotide of the incorrect allele (e.g., an ASO specific for a mutant allele cross-hybridizing to a wild-type allele). In more preferred embodiments of the present invention, there is at least a five-fold signal differential. In highly preferred embodiments of the present invention, there is at least an order of magnitude signal differential between the ASO hybridizing to a target polynucleotide of the correct allele and the level of the

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signal produced by the ASO cross-hybridizing to a target polynucleotide of the incorrect allele.

While certain methods for detecting polymorphisms are described herein, other detection methodologies may be utilized. For example, additional methodologies are known and set forth, in Birren et al., Genome Analysis, 4:135-186, A Laboratory Manual. Mapping Genomes, Cold Spring Harbor Laboratory Press, Cold Spring Harbor, NY (1999); Maliga et al., Methods in Plant Molecular Biology. A Laboratory Course Manual, Cold Spring Harbor Laboratory Press, Cold Spring Harbor, NY (1995); Paterson, Biotechnology Intelligence Unit: Genome Mapping in Plants, R.G. Landes Co., Georgetown, TX, and Academic Press, San Diego, CA (1996); The Maize Handbook, Freeling and Walbot, eds., Springer-Verlag, New York, NY (1994); Methods in Molecular Medicine: Molecular Diagnosis of Genetic Diseases, Elles, ed., Humana Press, Totowa, NJ (1996); Clark, ed., Plant Molecular Biology: A Laboratory Manual, Clark, ed., Springer-Verlag, Berlin, Germany (1997), all of which are herein incorporated by reference in their entirety.

Detection of one or more of the polymorphisms, preferably one or more of the characterized polymorphisms, may be carried out using a collection of nucleic acid markers.

Preferred aspects of this invention comprise collections of nucleic acid markers comprising nucleic acid molecules where the collections range in size from about 10 non-identical members or more, to at least about 100 or 270 or higher, more preferably at least about 300 or 350, most preferably at least 400 or 500 or higher, up to about 1,000, or 2000 or even higher, say about 4,000 or greater, or more non-identical members. As used herein a non-identical member is a member that differs in nucleic acid or amino acid sequence. For example, a non-identical nucleic acid molecule is a nucleic acid molecule that differs in nucleic acid sequence from the nucleic acid molecule to which it is being compared. For example a nucleic acid molecule having the sequence 5' CCC 3' is not

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identical – *i.e.* is non-identical – to a nucleic acid molecule having the sequence 5' CCG 3'. In one limited aspect a collection may comprise all of the nucleic acid markers identified by this invention. Collections of nucleic acid markers can be located or organized in a variety of forms, *e.g.* on microarrays, in solutions, in bacterial clone libraries, *etc.* As used herein, an "organized" collection is a collection where the nucleic acid or amino acid sequence of a member of such a collection can be determined based on its physical location.

In order to simultaneously screen for multiple polymorphisms, the nucleic acid markers can be designed for simultaneous use known as multiplexing. Examples of design approaches for multiplexing are set forth in Cho *et al.*, *Nature Genetics 23:*203-207 (1999); Wang *et al.*, *Science 280:* 1077-1082 (1998), the entirety of which is herein incorporated by reference; Winzeler *et al.*, *Curr. Opin. Genet. Dev. 4:* 602-608 (1997), the entirety of which is herein incorporated by reference. Examples of nucleic acid markers that have been optimized for multiplexing are the primers set forth in Table B. Multiplex parameters often require the selection of loci with similar amplification efficiencies, minimizing the concentration of the primers used, and an increased magnesium concentration (Cho *et al.*, *Nature Genetics 23:*203-207 (1999)).

In a preferred embodiment, the polymorphism is present and screened for in a mapping population, *e.g.* a collection of plants capable of being used with markers such as polymorphic markers to map genetic position of traits. The choice of appropriate mapping population often depends on the type of marker systems employed (Tanksley *et al.*, *J.P. Gustafson and R. Appels* (eds.). Plenum Press, New York, pp. 157-173 (1988), the entirety of which is herein incorporated by reference). Consideration must be given to the source of parents (adapted vs. exotic) used in the mapping population. Chromosome pairing and recombination rates can be severely disturbed (suppressed) in wide crosses (adapted x exotic) and generally yield greatly reduced linkage distances. Wide crosses

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will usually provide segregating populations with a relatively large number of polymorphisms when compared to progeny in a narrow cross (adapted x adapted).

An F<sub>2</sub> population is the first generation of selfing (self-pollinating) after the hybrid seed is produced. Usually a single F<sub>1</sub> plant is selfed to generate a population segregating for all the genes in Mendelian (1:2:1) pattern. Maximum genetic information is obtained from a completely classified F<sub>2</sub> population using a codominant marker system (Mather, Measurement of Linkage in Heredity: Methuen and Co., (1938), the entirety of which is herein incorporated by reference). In the case of dominant markers, progeny tests (*e.g.*, F<sub>3</sub>, BCF<sub>2</sub>) are required to identify the heterozygotes, in order to classify the population. However, this procedure is often prohibitive because of the cost and time involved in progeny testing. Progeny testing of F<sub>2</sub> individuals is often used in map construction where phenotypes do not consistently reflect genotype (*e.g.* disease resistance) or where trait expression is controlled by a QTL. Segregation data from progeny test populations *e.g.* F<sub>3</sub> or BCF<sub>2</sub>) can be used in map construction. Marker-assisted selection can then be applied to cross progeny based on marker-trait map associations (F<sub>2</sub>, F<sub>3</sub>), where linkage groups have not been completely disassociated by recombination events (*i.e.*, maximum disequilibrium).

Recombinant inbred lines (RIL) (genetically related lines; usually >F<sub>5</sub>, developed from continuously selfing F<sub>2</sub> lines towards homozygosity) can be used as a mapping population. Information obtained from dominant markers can be maximized by using RIL because all loci are homozygous or nearly so. Under conditions of tight linkage (*i.e.*, about <10% recombination), dominant and co-dominant markers evaluated in RIL populations provide more information per individual than either marker type in backcross populations (Reiter. *Proc. Natl. Acad. Sci. (U.S.A.)* 89:1477-1481 (1992), the entirety of which is herein incorporated by reference). However, as the distance between markers becomes larger (*i.e.*, loci become more independent), the information in RIL populations decreases dramatically when compared to codominant markers.

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Backcross populations (e.g., generated from a cross between a successful variety (recurrent parent) and another variety (donor parent) carrying a trait not present in the former) can be utilized as a mapping population. A series of backcrosses to the recurrent parent can be made to recover most of its desirable traits. Thus a population is created consisting of individuals nearly like the recurrent parent but each individual carries varying amounts or mosaic of genomic regions from the donor parent. Backcross populations can be useful for mapping dominant markers if all loci in the recurrent parent are homozygous and the donor and recurrent parent have contrasting polymorphic marker alleles (Reiter et al., Proc. Natl. Acad. Sci. (U.S.A.) 89:1477-1481 (1992), the entirety of which is herein incorporated by reference). Information obtained from backcross populations using either codominant or dominant markers is less than that obtained from F<sub>2</sub> populations because one, rather than two, recombinant gamete is sampled per plant. Backcross populations, however, are more informative (at low marker saturation) when compared to RILs as the distance between linked loci increases in RIL populations (i.e. about .15% recombination). Increased recombination can be beneficial for resolution of tight linkages, but may be undesirable in the construction of maps with low marker saturation.

Near-isogenic lines (NIL) (created by many backcrosses to produce a collection of individuals that is nearly identical in genetic composition except for the trait or genomic region under interrogation) can be used as a mapping population. In mapping with NILs, only a portion of the polymorphic loci is expected to map to a selected region.

Bulk segregant analysis (BSA) is a method developed for the rapid identification of linkage between markers and traits of interest (Michelmore *et al.*, *Proc. Natl. Acad. Sci. U.S.A.* 88:9828-9832 (1991), the entirety of which is herein incorporated by reference). In BSA, two bulked DNA samples are drawn from a segregating population originating from a single cross. These bulks contain individuals that are identical for a particular trait (resistant or susceptible to particular disease) or genomic region but

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arbitrary at unlinked regions (i.e. heterozygous). Regions unlinked to the target region will not differ between the bulked samples of many individuals in BSA.

While any appropriate mapping population may be used in conjunction with this invention, in a preferred embodiment the mapping population is an *Arabidopsis* population, where the population was created, at least in part, by crossing two different *Arabidopsis* ecotypes, where one of the ecotypes has a phenotype of interest. In an even more preferred embodiment the ecotypes are *Arabidopsis*, *thaliana*, Columbia and *Arabidopsis*, *thaliana*, Landsberg *erecta*. In another preferred embodiment, the mapping population is an *Arabidopsis* population, where the population was created, at least in part, by crossing two different *Arabidopsis* ecotypes, where one of the ecotypes has a phenotype of interest, propagating and self pollinating seeds from such a cross and selecting a collection of plants with the phenotype of interest to be the mapping population.

Classical mapping studies often utilize easily observable, visible traits instead of molecular markers. These visible traits are also known as naked eye polymorphisms. These traits can be morphological like plant height, fruit size, shape and color or physiological like disease response, photoperiod sensitivity and crop maturity. Visible traits are useful and are still in use because they represent actual phenotypes and are easy to score without any specialized lab equipment. By contrast, many nucleic acid markers are arbitrary loci for use in linkage mapping and often not associated with specific plant phenotypes (Young, *Encyclopedia of Agricultural Science*, Vol. 3, pp. 275-282 (1994), the entirety of which is herein incorporated by reference). Many morphological markers cause such large effects on phenotype that they are undesirable in breeding programs. Many other visible traits have the disadvantage of being developmentally regulated (*i.e.*, expressed only at certain stages; or in specific tissue and organs). Oftentimes, visible traits mask the effects of linked minor genes making it nearly impossible to identify

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desirable linkages for selection (Tanksley et al., Biotech. 7:257-264 (1989), the entirety of which is herein incorporated by reference).

Although a number of important agronomic characteristics are controlled by loci having major effects on phenotype, many economically important traits, such as yield and some forms of disease resistance, are quantitative in nature. This type of phenotypic variation in a trait is typically characterized by continuous, normal distribution of phenotypic values in a particular population (polygenic traits) (Beckmann and Soller, *Oxford Surveys of Plant Molecular Biology, Miffen.* (ed.), Vol. 3, Oxford University Press, UK., pp. 196-250 (1986), the entirety of which is herein incorporated by reference). Loci contributing to such genetic variation are often termed minor genes, as opposed to major genes with large effects that follow a Mendelian pattern of inheritance. Polygenic traits are also predicted to follow a Mendelian type of inheritance, however the contribution of each locus is expressed as an increase or decrease in the final trait value. The nucleic acid markers of the present invention can be used to identify and isolate nucleic acid regions or molecules associated with desired polygenic or single gene traits.

In one embodiment, the nucleic acid markers of the present invention are used to isolate or identify an allele of a quantitative trait locus or Mendelian locus.

Nucleic acid markers of the present invention capable of detecting one or more of the polymorphisms may be employed in genetic or physical studies using linkage analysis. Mapping marker genetic locations is based on the observation that two markers located near each other on the same chromosome will tend to be passed together from parent to offspring. During gamete production, DNA strands occasionally break and rejoin in different places on the same chromosome or on the homologous chromosome. The closer the markers are to each other, the more tightly linked and the less likely a recombination event will fall between and separate them. Recombination frequency thus provides an estimate of the distance between two markers.

Linkage analysis is based on the level at which markers and genes are co-inherited (Rothwell, *Understanding Genetics. 4<sup>th</sup> Ed.* Oxford University Press, New York, p. 703 (1988), the entirety of which is herein incorporated by reference). Statistical tests like chi-square analysis can be used to test the randomness of segregation or linkage (Kochert, *The Rockefeller Foundation International Program on Rice Biotechnology*, University of Georgia Athens, GA, pp. 1-14 (1989), the entirety of which is herein incorporated by reference). In linkage mapping, the proportion of recombinant individuals out of the total mapping population provides the information for determining the genetic distance between the loci (Young, *Encyclopedia of Agricultural Science, Vol. 3*, pp. 275-282 (1994), the entirety of which is herein incorporated by reference). Any statistical analysis that establishes linkage may be used. An example of a suitable linkage approach is Intermap as set forth in Cho *et al.*, *Nature Genetics 23*: 203-207 (1999). Example 6 sets forth another exemplary linkage approach.

In segregating populations, target genes have been reported to have been placed within an interval of 5-10 cM with a high degree of certainty (Tanksley *et al.*, *Trends in Genetics 11(2):63-68* (1995), the entirety of which is herein incorporated by reference). The markers defining this interval are used to screen a larger segregating population to identify individuals derived from one or more gametes containing a crossover in the given interval. Such individuals are useful in orienting other markers closer to the target gene. Once identified, these individuals can be analyzed in relation to all molecular markers within the region to identify those closest to the target.

Markers of the present invention can be employed to locate genes. The genetic linkage of additional marker molecules can be established by a genetic mapping model such as, without limitation, the flanking marker model reported by Lander and Botstein, *Genetics 121*:185-199 (1989), the entirety of which is herein incorporated by reference, and the interval mapping, based on maximum likelihood methods described by Lander and Botstein, *Genetics 121*:185-199 (1989), the entirety of which is herein incorporated

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by reference and implemented in the software package MAPMAKER/QTL (Lincoln and Lander, *Mapping Genes Controlling Quantitative Traits Using MAPMAKER/QTL*, Whitehead Institute for Biomedical Research, Massachusetts, (1990), the entirety of which is herein incorporated by reference). Additional software includes Qgene, Version 2.23 (Department of Plant Breeding and Biometry, 266 Emerson Hall, Cornell University, Ithaca, NY (1996), the manual of which is herein incorporated by reference in its entirety).

The LOD score essentially indicates how much more likely the data are to have arisen assuming the presence of an allele than in its absence. The LOD threshold value for avoiding a false positive with a given confidence, say 95%, depends on the number of markers and the length of the genome. Graphs indicating LOD thresholds are set forth in Lander and Botstein, *Genetics 121*:185-199 (1989), the entirety of which is herein incorporated by reference and further described by Arús and Moreno-González, *Plant Breeding*, Hayward, Bosemark, Romagosa (eds.) Chapman & Hall, London, pp. 314-331 (1993), the entirety of which is herein incorporated by reference.

In a preferred embodiment of the present invention the nucleic acid marker exhibits a LOD score of greater than 2.0, more preferably 2.5, even more preferably greater than 3.0 or 4.0 with the trait or phenotype of interest.

Additional models can be used. Many modifications and alternative approaches to interval mapping have been reported, including the use of non-parametric methods (Kruglyak and Lander, *Genetics*, 139:1421-1428 (1995), the entirety of which is herein incorporated by reference). Multiple regression methods or models can be also used, in which the trait is regressed on a large number of markers (Jansen, *Biometrics in Plant Breed*, van Oijen, Jansen (eds.) Proceedings of the Ninth Meeting of the Eucarpia Section Biometrics in Plant Breeding, The Netherlands, pp. 116-124 (1994); Weber and Wricke, *Advances in Plant Breeding*, Blackwell, Berlin, 16 (1994), the entirety of which is herein incorporated by reference). Procedures combining interval mapping with regression

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analysis, whereby the phenotype is regressed onto a single putative QTL at a given interval, and at the same time onto a number of polymorphisms that serve as 'cofactors,' have been reported by Jansen and Stam, *Genetics*, 136:1447-1455 (1994), the entirety of which is herein incorporated by reference and Zeng, *Genetics*, 136:1457-1468 (1994), the entirety of which is herein incorporated by reference. Generally, the use of cofactors reduces the bias and sampling error of the estimated QTL positions (Utz and Melchinger, *Biometrics in Plant Breeding*, van Oijen, Jansen (eds.) Proceedings of the Ninth Meeting of the Eucarpia Section Biometrics in Plant Breeding, The Netherlands, pp.195-204 (1994)), thereby improving the precision and efficiency of QTL mapping (Zeng, *Genetics*, 136:1457-1468 (1994), the entirety of which is herein incorporated by reference). These models can be extended to multi-environment experiments to analyze genotype-environment interactions (Jansen *et al.*, *Theo. Appl. Genet. 91:33-37* (1995), the entirety of which is herein incorporated by reference).

The nucleic acid markers of the present invention may be used to isolate an allele, a region of genomic DNA associated with a phenotype, *etc*. Once the genomic region associated with the phenotype of interest is defined relative to at least one nucleic acid marker, preferably at least two nucleic acid markers capable of detecting different polymorphisms, the genomic region associated with the phenotype may be further characterized. One approach is to select additional nucleic acid markers from the genomic region associated with the trait and localize the genomic region associated with the phenotype to a smaller genomic region by a technique such as fine mapping.

For example, in a preferred embodiment a method for identifying or isolating a genomic region associated with a phenotypic trait that comprises (A) screening a mapping population of *Arabidopsis* plants to determine the linkage of the phenotypic trait with a first collection of polymorphisms, wherein the first collection of polymorphisms is distributed throughout the genome of the mapping population of *Arabidopsis* plants at an average density of more than one polymorphism per about 500kb - 100kb; (B) calculating

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the linkage of each of the first collection of polymorphisms to the phenotypic trait;

(C) identifying a genomic region most closely associated with the phenotypic trait;

(D) selecting a second collection of polymorphisms from the genomic region; and

(E) screening the mapping population of *Arabidopsis* plants to determine the linkage of the phenotypic trait with the second collection of polymorphisms from the genomic region, wherein the second collection of polymorphisms have an average density of more than one polymorphism per about 50kb - 1kb.

In an embodiment of the present invention, for a fine mapping step of the present invention the collection of marker nucleic acids is capable of detecting a characterized polymorphism at a density of greater than one polymorphism per 50kb, more preferably at a density greater than one polymorphism per 25kb, even more preferably at a density greater than one polymorphism per 10kb or 5kb. It is understood, that the fine mapping using such a collection of markers may be carried out, for example, in a single assay or simultaneously.

15 Once the genomic region associated with the phenotype is identified, the genomic region may be isolated. Alternatively, or in conjunction, such a region may be further defined or characterized. Many approaches are known in the art and may be undertaken (Sambrook et al., Molecular Cloning 1: A Laboratory Manual, 2d ed., Ford et al., eds., Cold Spring Harbor Laboratory Press, Cold Spring Harbor, NY (1989); Sambrook et al., Molecular Cloning 2: A Laboratory Manual, 2d ed., Ford et al., eds., Cold Spring 20 Harbor Laboratory Press, Cold Spring Harbor, NY (1989); Sambrook et al., Molecular Cloning 3: A Laboratory Manual, 2d ed., Ford et al., eds., Cold Spring Harbor Laboratory Press, Cold Spring Harbor, NY (1989); Maliga et al., Methods in Plant Molecular Biology: A Laboratory Course Manual, Cold Spring Harbor Laboratory Press, 25 Cold Spring Harbor, NY (1995); and Birren et al., Genome Analysis: A Laboratory Manual. Volume 2: Detecting Genes, Cold Spring Harbor Laboratory Press, Cold Spring Harbor, NY (1998), all of which are herein incorporated by reference in their entirety).

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phenotype may be determined and subjected to bioinformatic analysis (Coulson, *Trends in Biotechnology 12:*76-80 (1994); Birren *et al.*, *Genome Analysis 1*, Cold Spring Harbor Laboratory Press, Cold Spring Harbor, New York 543-559 (1997); Huang, *et al.*, *Genomics 46:*37-45 (1997), all of which are herein incorporated by reference in their entirety). Such bioinformatic approaches can provide, for example, information on the location of putative open reading frames, promoters, and a variety of nucleotide motifs. Moreover, also using bioinformatic approaches, the nucleic acid sequence of the genomic region can be compared with other nucleic acid sequences. Such comparisons can

For example, once identified, the sequence of the genomic region associated with the

Other methods can be utilized to further isolate, define, or characterize the genomic region associated with the phenotype. The expression profiles of mRNA and proteins derived from genes that are located within the genetic region associated with the phenotype can be analyzed. Such analysis, will in certain circumstances, allow the gene or genes associated with the phenotype to be determined.

facilitate the isolation of Arabidopsis homologs to known genes or genomic regions.

Examples of such bioinformation tools are BLAST, GeneScan, GeneMark and AAT.

A genomic region or sub-region thereof may be isolated using any of the many techniques in the art. In addition to those procedures and methods set forth herein, practitioners are familiar with the standard resource materials which describe specific conditions and procedures for the construction, manipulation and isolation of macromolecules (e.g., DNA molecules, plasmids, etc.), generation of recombinant organisms and the screening and isolating of clones, (see, for example, Sambrook et al., Molecular Cloning: A Laboratory Manual, Cold Spring Harbor Press (1989); Mailga et al., Methods in Plant Molecular Biology, Cold Spring Harbor Press (1995); Birren et al., Genome Analysis: Analyzing DNA, 1, Cold Spring Harbor, New York, all of which are herein incorporated by reference in their entirety).

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The biological function of a genomic region or subregion thereof such as a gene or open reading frame, can be further investigated using a mutant complementation approach or other reverse genetics approach. For example, a gene or genes identified within the genomic region associated with the phenotype may be isolated from the organism exhibiting the non-mutant phenotype (often referred to as the wild type). Such a gene or genes may be introduced into an appropriate organism that lacks the phenotype (often referred to as mutant) either by crosses or by molecular genetic techniques such as transformation or transfection. Organisms having the introduced genetic material may be screened to determine whether the introduced gene or genes complements, i.e. restores the phenotype of the mutant (Pan, FEBS Lett. 459(3): 405-410 (1999); Kerckhoffs et al., Mol. Gen. Genet. 6: 901-907 (1999); Lizotte et al., Gene 234(1): 35-44 (1999); Berna et al., Genetics 152: 729-742 (1999); Liu et al., Proc. Natl. Acad. Sci. (USA) 96(11): 6535-6540 (1999); Pia et al., Plant Physiol. 119(4): 1527-1534 (1999); Loulergue et al., Gene 225(1-2): 47-57 (1998); Jouannic et al., Eur. J. Bioche, 258(2): 402-410 (1998), all of which are herein incorporated by reference in their entirety). While gene or genes etc. may be introduced into any organism, preferred organisms are plants, yeasts, and bacteria particularly E. coli. In a more preferred embodiment the organism is Arabidopsis.

The nucleic acid markers of the present invention may be used for chromosomal walking. Such walking, in conjunction with linkage analysis, can enable the isolation of genes. Once a nucleic acid marker is linked to a region of interest, the chromosome walking technique can be used to find the genes via overlapping clones. For chromosome walking, random molecular markers or established molecular linkage maps are used to conduct a search to localize the gene adjacent to one or more markers capable of detecting a polymorphism. A chromosome walk (Bukanov and Berg, *Mo. Microbiol.* 11:509-523 (1994), the entirety of which is herein incorporated by reference; Birkenbihl and Vielmetter, *Nucleic Acids Res.* 17:5057-5069 (1989), the entirety of which is herein incorporated by reference; Wenzel and Herrmann, *Nucleic Acids Res.* 16:8323-8336,

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(1988), the entirety of which is herein incorporated by reference) is then initiated from the closest linked marker. Starting from the selected clones, labeled probes specific for the ends of the insert DNA are synthesized and used as probes in hybridizations against a representative library. Clones hybridizing with one of the probes are picked and serve as templates for the synthesis of new probes; by subsequent analysis, contigs are produced.

The degree of overlap of the hybridizing clones used to produce a contig can be determined by comparative restriction analysis. Comparative restriction analysis can be carried out in different ways all of which exploit the same principle; two clones of a library are very likely to overlap if they contain a limited number of restriction sites for one or more restriction endonucleases located at the same distance from each other. The most frequently used procedures are, fingerprinting (Coulson *et al.*, *Proc. Natl. Acad. Sci.* (U.S.A.) 83:7821-7821, (1986), the entirety of which is herein incorporated by reference; Knott *et al.*, *Nucleic Acids Res. 16*:2601-2612 (1988), the entirety of which is herein incorporated by reference; Eiglmeier *et al. Mol. Microbiol.* 7:197-206 (1993), the entirety of which is herein incorporated by reference), restriction fragment mapping (Smith and Birnstiel, *Nucleic Acids Res.* 3:2387-2398 (1976), the entirety of which is herein incorporated by reference), and the "landmarking" technique (Charlebois *et al. J. Mol. Biol.* 222:509-524 (1991), the entirety of which is herein incorporated by reference).

It is understood that the nucleic acid molecules of the present invention may in one embodiment be used for chromosomal walking. In a preferred embodiment, nucleic acid molecules of the present invention may in one embodiment be used in the chromosomal walking of *Brassicaceae*, particularly *Arabidopsis*.

Nucleic acid markers of the present invention can be used in comparative mapping and comparative chromosomal walking. Comparative mapping within families provides a method to assess the degree of sequence conservation, gene order, ploidy of species, ancestral relationships and the rates at which individual genomes are evolving. It also provides a method to isolate genetic regions or sub-aspects thereof such as genes.

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Comparative mapping has been carried out by utilizing molecular markers from one species with another species. As in genetic mapping, nucleic acid markers are needed but instead of direct hybridization to mapping filters, the markers can also be used to select large insert clones from a total genomic DNA library of a related species. The selected clones can then be used to physically map the region in the target species. The advantage of this method for comparative mapping is that no mapping population or linkage map of the target species is needed and the clones may also be used in other closely related species. By comparing the results obtained by genetic mapping in model plants, with those from other species, similarities of genomic structure among plants species can be established. Comparative mapping using nucleic acid markers of the present invention permits the identification and/or isolation of non-*Arabidopsis* syntenic regions and homolog genes with such regions.

It is understood that nucleic acid markers of the present invention may in another embodiment be used in comparative mapping. In a preferred embodiment the markers of the present invention may be used in the comparative mapping of non-*Arabidopsis* plant species, including but not limited to alfalfa, barley, *Brassica*, broccoli, cabbage, citrus, cotton, garlic, oat, oilseed rape, onion, canola, flax, an ornamental plant, maize, pea, peanut, pepper, potato, rice, rye, sorghum, soybean, strawberry, sugarcane, sugarbeet, tomato, wheat, poplar, pine, fir, eucalyptus, apple, lettuce, lentils, grape, banana, tea, turf grasses, sunflower, oil palm, *Phaseolus etc*. Particularly preferred non-*Arabidopsis* plants to utilize for comparative mapping are the *Brassicaceae*.

Agents of the present invention include nucleic acid molecules and more specifically include nucleic acid markers capable of detecting polymorphisms. In a preferred embodiment the nucleic acid molecules of the present invention are derived from *Arabidopsis* and in an even more preferred embodiment the nucleic acid molecules of the present invention are derived from *Arabidopsis thaliana*, Landsberg *erecta* or *Arabidopsis thaliana*, Columbia.

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In another preferred embodiment, the nucleic acid molecules of the present invention include those isolated utilizing the nucleic acid markers of the present invention. The present invention also encompasses the use of these and other nucleic acids of the present invention in recombinant constructs. Using methods known to those of ordinary skill in the art, such molecules can be introduced into a host cell or organism of choice. Potential host cells include both prokaryotic and eukaryotic cells. A host cell may be unicellular or found in a multicellular differentiated or undifferentiated organism depending upon the intended use. It is understood that useful exogenous genetic material may be introduced into any cell or organism such as a plant cell, plant, mammalian cell, mammal, fish cell, fish, bird cell, bird or bacterial cell.

In a preferred embodiment the exogenous DNA is introduced into a plant in a suitable construct. Preferred plants are selected from the group consisting of: alfalfa, *Arabidopsis*, barley, *Brassica*, broccoli, cabbage, citrus, cotton, garlic, oat, oilseed rape, onion, canola, flax, an ornamental plant, peanut, pepper, potato, rice, rye, sorghum, strawberry, sugarcane, sugarbeet, tomato, wheat, poplar, pine, fir, eucalyptus, apple, lettuce, lentils, grape, banana, tea, turf grasses, sunflower, soybean, and *Phaseolus*. A particularly preferred group of plants is rice, cotton, wheat, maize and soybean.

As used herein, an agent, be it a naturally occurring molecule or otherwise may be "substantially purified," if, referring to a molecule separated from substantially all other molecules normally associated with it in its native state. More preferably a substantially purified molecule is the predominant species present in a preparation. A substantially purified molecule may be greater than 60% free, preferably 75% free, more preferably 90% free, and most preferably 95% free from the other molecules (exclusive of solvent) present in the natural mixture. The term "substantially purified" is not intended to encompass molecules present in their native state.

The agents of the present invention will preferably be "biologically active" with respect to either a structural attribute, such as the capacity of a nucleic acid to hybridize to

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another nucleic acid molecule, or the ability of a protein to be bound by an antibody (or to compete with another molecule for such binding). Alternatively, such an attribute may be catalytic, and thus involve the capacity of the agent to mediate a chemical reaction or response.

The agents of the present invention may can also be recombinant. As used herein, the term recombinant describes (a) nucleic acid molecules that are constructed or modified outside of cells and that can replicate or function in a living cell, (b) molecules that result from the transcription, replication or translation of recombinant nucleic acid molecules , or (c) organisms that contain recombinant nucleic acid molecules or are modified using recombinant nucleic acid molecules.

It is understood that the agents of the present invention may be labeled with reagents that facilitate detection of the agent (e.g. fluorescent labels, Prober et al., Science 238:336-340 (1987); Albarella et al., EP 144914, chemical labels, Sheldon et al., U.S. Patent 4,582,789; Albarella et al., U.S. Patent 4,563,417, modified bases, Miyoshi et al., EP 119448, all of which are herein incorporated by reference in their entirety).

Fragment nucleic acid molecules may encode significant portion(s) of, or indeed most of, these nucleic acid molecules. For example, a fragment nucleic acid molecule can encode an *Arabidopsis* protein or fragment thereof. Alternatively, the fragments may comprise smaller oligonucleotides. Exemplary fragment sizes include fragments having from about 15 to about 400 nucleotide residues and more preferably, about 15 to about 30 nucleotide residues, or about 50 to about 100 nucleotide residues, or about 100 to about 200 nucleotide residues, or about 275 to about 350 nucleotide residues.

Nucleic acid molecules or fragments thereof of the present invention are capable of specifically hybridizing to other nucleic acid molecules under certain circumstances.

As used herein, two nucleic acid molecules are said to be capable of specifically hybridizing to one another if the two molecules are capable of forming an anti-parallel,

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double-stranded nucleic acid structure. A nucleic acid molecule is said to be the "complement" of another nucleic acid molecule if they exhibit complete complementarity. As used herein, molecules are said to exhibit "complete complementarity" when every nucleotide of one of the molecules is complementary to a nucleotide of the other. Two molecules are said to be "minimally complementary" if they can hybridize to one another with sufficient stability to permit them to remain annealed to one another under at least conventional "low-stringency" conditions. Similarly, the molecules are said to be "complementary" if they can hybridize to one another with sufficient stability to permit them to remain annealed to one another under conventional "high-stringency" conditions. Conventional stringency conditions are described by Sambrook et al., Molecular Cloning, A Laboratory Manual, 2nd Ed., Cold Spring Harbor Press, Cold Spring Harbor, New York (1989), and by Haymes et al. Nucleic Acid Hybridization, A Practical Approach, IRL Press, Washington, DC (1985), the entirety of which is herein incorporated by reference. Departures from complete complementarity are therefore permissible, as long as such departures do not completely preclude the capacity of the molecules to form a double-stranded structure. Thus, in order for a nucleic acid molecule to serve as a primer or probe it need only be sufficiently complementary in sequence to be able to form a stable double-stranded structure under the particular solvent and salt concentrations employed.

Appropriate stringency conditions which promote DNA hybridization, for example, 6.0 X sodium chloride/sodium citrate (SSC) at about 45°C, followed by a wash of 2.0 X SSC at 50°C, are known to those skilled in the art or can be found in *Current Protocols in Molecular Biology*, John Wiley & Sons, N.Y. (1989), 6.3.1-6.3.6, the entirety of which is herein incorporated by reference. For example, the salt concentration in the wash step can be selected from a low stringency of about 2.0 X SSC at 50°C to a high stringency of about 0.2 X SSC at 50°C. In addition, the temperature in the wash step can be increased from low stringency conditions at room temperature, about 22°C, to high

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stringency conditions at about 65°C. Both temperature and salt may be varied, or either the temperature or the salt concentration may be held constant while the other variable is changed.

Hybridizations involving at least one oligonucleotide can necessitate changes from the above hybridization conditions. Highly stringent conditions are often selected to be equal to the  $T_m$  point for a particular probe. Sometimes the term " $T_d$ " is used to define the temperature at which at least half of the probe dissociates from a perfectly matched target nucleic acid. In any case, a variety of estimation techniques for estimating the  $T_m$  or  $T_d$  are available, and generally described in Tijssen, id. Typically, G-C base pairs in a duplex are estimated to contribute about 3°C to the  $T_m$ , while A-T base pairs are estimated to contribute about 2°C, up to a theoretical maximum of about 80-100°C. However, more sophisticated models of  $T_M$  and  $T_d$  are available and appropriate in which G-C stacking interactions, solvent effects, the desired assay temperature and the like are taken into account. For example, PCR primers can be designed to have a dissociation temperature ( $T_d$ ) of approximately 60°C, using the formula:  $T_d = ((((3 \times \#GC) + (2 \times \#AT)) \times 37) - 562) / \#bp) - 5$ ; where #GC, #AT, and #bp are the number of guanine-cytosine base pairs, the number of adenine-thymine base pairs, and the number of total base pairs, respectively, involved in the annealing of the primer to the template DNA.

Nucleic acid markers of the present invention can be used to characterize transformants or germplasm, as a genetic diagnostic test for plant breeding or to identify individuals or varieties (Soller and Beckmann, *Theor. Appl. Genet.* (67):25-33 (1983), the entirety of which is herein incorporated by reference). Such markers can also be used to obtain information about: (1) the number, effect, and chromosomal location of each gene affecting a trait; (2) effects of multiple copies of individual genes (gene dosage); (3) interaction between/among genes controlling a trait (epistasis); (4) whether individual genes affect more than one trait (pleiotropy); and (5) stability of gene function across environments (Gx E interactions).

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In a preferred embodiment, the nucleic acid markers of the present invention may be used in marker assisted introgression of traits into plants. Marker assisted introgression involves the transfer of a chromosome region defined by one or more markers from one germplasm to a second germplasm. An initial step in such a process is the localization of the trait or region by mapping. One use of marker assisted introgression of genomic regions is in the generation of near isogenic lines (NILs) or recombinant near isogenic lines (RILs). In one aspect of the present invention, the nucleic acid markers are used to generate *Arabidopsis* NILs or RILs. As used herein, introgression is the process of transferring a genetic region from one genetic background to a second but non-identical genetic background.

Additional markers, such as AFLP markers, RFLP markers, RAPD markers, SNPs, phenotypic markers, isozyme markers can be utilized in combination with or separately from the markers of the invention (Walton, Seed World 22-29 (1993), the entirety of which is herein incorporated by reference; Burow and Blake, *Molecular Dissection of Complex Traits*, 13-29, Eds. Paterson, CRC Press, New York (1988), the entirety of which is herein incorporated by reference). Examples of additional markers are set forth in Cho *et al.*, *Nature Genetics* 23: 203-205 (1999).

DNA markers can be developed from nucleic acid molecules using restriction endonucleases, the PCR and/or DNA sequence information. RFLP can result from single base changes or insertions/deletions. RFLP are highly abundant in plant genomes, have a medium level of polymorphism and are developed by a combination of restriction endonuclease digestion and Southern blotting hybridization. CAPS are similarly developed from restriction nuclease digestion but only of specific PCR products. CAPS are also codominant, have a medium level of polymorphism and are highly abundant in the genome. The CAPS result from single base changes and insertions/deletions. RAPDs are developed from DNA amplification with random primers and result from single base changes and insertions/deletions in plant genomes. RAPDs with a medium level of

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polymorphisms are highly abundant. AFLP markers require using the PCR on a subset of restriction fragments from extended adapter primers. AFLPs are both dominant and codominant are highly abundant in genomes and exhibit a medium level of polymorphism. SSRs require DNA sequence information. SSRs result from repeat length changes, are highly polymorphic, and do not exhibit as high a degree of abundance in the genome as CAPS, AFLPs and RAPDs. SNPs also require DNA sequence information. SNPs result from single base substitutions. They are highly abundant and exhibit a medium of polymorphism (Rafalski *et al.*, In: *Nonmammalian Genomic Analysis*, ed. Birren and Lai, Academic Press, San Diego, CA, pp. 75-134 (1996), the entirety of which is herein incorporated by reference).

### Computer Readable Media

A polymorphism or nucleic acid molecule of the present invention can be "provided" in a variety of mediums to facilitate use. Moreover, the nucleic acid markers and other nucleic acid molecules of the present invention may also be so presented.

In one embodiment, a polymorphism may be presented in a manner that sets forth 1, more preferably 2, 3, 4, 5, 6, or 7 of the following features alone or in combination with other features: (1) type of polymorphism (e.g. SNP, insertion, deletion etc.); (2) physical location of the polymorphism on a chromosome; (3) nucleotide sequence variation associated with one or more of the alleles; (4) nucleotide sequences of nucleic acid marker molecules capable of detecting the polymorphism; (5) physical location of the polymorphism relative to a piece of isolated DNA (e.g., BAC); (6) methodology for detecting the polymorphism; (7) physical distance from that polymorphism to another polymorphism; and (8) genetic linkage with a phenotype or other polymorphism.

Such a medium can also provide a subset thereof in a form that allows a skilled artisan to examine these features.

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In one application of this embodiment, a polymorphism and associated features of the present invention can be recorded on computer readable media. In another embodiment, a nucleic acid sequence of the present invention can be recorded on computer readable media alone or in combination with a polymorphisms and associated features. As used herein, "computer readable media" refers to any medium that can be read or accessed by a computer, either directly or indirectly through a network. Such media include, but are not limited to: magnetic storage media, such as disks or magnetic tape; optical storage media such as optical disks; electrical storage media such as read-only memory (ROM) or Random Access Memory (RAM); and hybrids of these categories such as magnetic/optical storage media. A skilled artisan can readily appreciate how any of the known computer readable mediums can be used to create a manufacture comprising computer readable medium having recorded thereon a nucleotide sequence of the present invention.

As used herein, "recorded" refers to a process for storing information on computer readable medium. A skilled artisan can readily adopt any of the known methods for recording information on computer readable medium to generate media comprising the information of the present invention. A variety of data storage structures are available to a skilled artisan for creating a computer readable medium having recorded thereon a nucleotide sequence of the present invention. The choice of the data storage structure will generally be based on the means chosen to access the stored information. In addition, a variety of application programs and formats can be used to store the information of the present invention on computer readable medium. The sequence information can be represented, for example, in a word processing file, formatted in commercially-available software such as WordPerfect and Microsoft Word, in a network-accessible format, such as an HTML file or web page, an ASCII file, or stored in a database application, such as DB2, Excel, Sybase, Oracle, or the like. A skilled artisan can readily adapt any number of data file formats (e.g., text file or database) or data

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structures in order to obtain computer readable medium having recorded thereon the information of the present invention.

A skilled artisan is provided with access to the information for a variety of purposes. Publicly available computer software allows a skilled artisan to access, for example, sequence information provided in a computer readable medium.

The present invention further provides systems, particularly computer-based systems, which contain the information described herein. As used herein, "a computer-based system" refers to the hardware, software, and data storage used to analyze the information including the nucleic acid sequence information of the present invention. The minimum hardware of the computer-based systems of the present invention comprises a central processing unit (CPU), input/output apparatus, and data storage. A skilled artisan can readily appreciate that any one of the currently available computer-based systems are suitable for use in the present invention.

As indicated above, the computer-based systems of the present invention comprise a data storage having stored therein a polymorphism and any associated information of the present invention and the necessary hardware and software for supporting and implementing a search. As used herein, "data storage" refers to memory that can store information of the present invention, or a memory access apparatus (hardware and/or software) that can access manufactures having recorded thereon the information of the present invention.

Having now generally described the invention, the same will be more readily understood through reference to the following examples which are provided by way of illustration, and are not intended to be limiting of the present invention, unless specified.

#### Example 1

Assembled *Arabidopsis thaliana*, Landsberg *erecta* nucleic acid sequence is generated essentially as set forth below:

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# **DNA Preparation**

DNA from *Arabidopsis thaliana*, Landsberg *erecta* seedlings is prepared by a CTAB genomic DNA isolation protocol as described by Dean *et al. Plant J* 2:69-81(1992) and modified by Dubois *et al. Plant J.* 13:141-151 (1998), the entirety of which is herein incorporated by reference.

A solution of DNA to be sheared is prepared in a 1.5 ml microcentrifuge tube by mixing 15  $\mu$ g of DNA, 6  $\mu$ l of 10X mung bean (MB) buffer (10X MB buffer = 300mM NaOAc, pH 5.0, 500 mM NaCl, 10 mM ZnCl<sub>2</sub>, 50% glycerol), and water to a final volume of 60  $\mu$ l. The DNA solution is kept on ice prior to sonication. For sonication, a cup horn probe chilled with ice water for 1 hour prior to sonication is used. The sonicator (Ultrasonic Liquid Processor XL2020 , Misonix Inc.) is pulsed for approximately 10 seconds on full power prior to use. DNA samples are sonicated twice for 6 seconds each at 60% power. Four sample tubes may be processed at once in a multi-tube rack which is positioned 1 to 3 mm above the opening in the probe. The DNA is returned to ice and a 1  $\mu$ l sample is analyzed by electrophoresis on a 0.8% agarose gel in 0.5X TBE gel, run at 60 volts for 30 minutes. Sonication may be repeated if necessary.

A 0.26  $\mu$ l aliquot of mung bean nuclease (150,000 u/ml) is added to sheared DNA and the sample is incubated at 30° C for 10 minutes. To stop the digestion, 20  $\mu$ l of 1 M NaCl, 140  $\mu$ l dd H<sub>2</sub>0, and 200 ml of phenol:chloroform are added to the sample which is then vortexed and centrifuged for 20 minutes at 13,000 rpm. The resulting aqueous phase is transferred into a new 1.5 ml microcentrifuge tube, 500  $\mu$ l of 95% ethanol is added, and the DNA is precipitated overnight at -80° C. The sample is centrifuged for 30 minutes at 13,000 rpm, washed with 500  $\mu$ l of 95% ethanol and centrifuged again for 30 minutes at 13,000 rpm. The sample is then dried under vacuum, and resuspended in 10  $\mu$ l TE.

The sheared DNA fragments are sized and purified by preparative agarose gel electrophoresis. Five microliters of 6x BP-XC-glycerol dye (0.25% BP, 0.25% XC, 30% glycerol) is added to the sample. The sample is split into two samples and loaded (12.5

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 $\mu$ l per lane) on a 0.8% (1x TAE) low-melting agarose gel (SeaPlaque GTG) and electrophoresed at 60 V, 46 mA for 3.5 hours.

The gel is photographed under long wave UV and slices containing DNA fragments of 1.3 - 1.7 kb and 2 - 4 kb are excised and excess agarose cut away. The gel slices are placed in 1.5 ml microcentrifuge tubes. One gel slice is stored at - $20^{\circ}$  C. 15  $\mu$ l of 1 M NaCl is added to the other gel slice, followed by melting of the agarose by incubation at  $65^{\circ}$  C for 8 minutes. The resulting approximately 250  $\mu$ l samples are placed into microcentrifuge tubes. An equal volume of water is added, following which the sample is vortexed and placed at room temperature for 2 minutes to bring the temperature up to 30 - $35^{\circ}$  C. 0.5 ml of water-saturated phenol that has been cooled on ice is added and the sample vortexed vigorously. The sample is placed on ice for 5 minutes, and the vortexing step repeated.

The sample is centrifuged at 4°C in a microcentrifuge for 20 minutes. The upper phase is transferred to a clean tube, and the bottom phenol layer is reextracted by addition of 200 µl of dd H<sub>2</sub>O. The sample is vortexed and placed on ice for 5 minutes, followed by centrifugation for 15 minutes. The aqueous layer is extracted and added to the aqueous layer from the previous step. Phenol extraction is repeated with 0.5 ml phenol, followed by vortexing and centrifugation for 20 minutes at 4°C. The aqueous layer is removed and repeated sec-butanol extractions are performed until the final volume is reduced to approximately 0.165 ml.

Two volumes of 95% ethanol (400  $\mu$ l) are added and the sample is stored at -80° C overnight. The sample is centrifuged for 30 minutes at room temperature to pellet the DNA, washed once with 95% ethanol and dried briefly under vacuum. The sample is resuspended in 7  $\mu$ l of TE. A 1  $\mu$ l sample is run on a 0.8% agarose gel with markers to estimate concentration of recovered fraction.

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#### M13 Library

20 ng of M13 DNA digested with SmaI is mixed with 1  $\mu$ l of 10x ligation buffer (10X ligation buffer = 0.5M tris pH 7.4, 0.1M MgCl<sub>2</sub>, 0.1M DDT), 1 $\mu$ l of 1mM ATP and 100 - 200 ng of sheared genomic DNA fragments (1 - 3  $\mu$ l volume), and 0.3  $\mu$ l of high concentration NEB ligase (5 unit/ $\mu$ l) is added. Water is added to a final volume of 10 $\mu$ l and the sample is incubated overnight at 14° C.

## Plasmid Library

200 ng (4  $\mu$ l) of pSTBlue vector (Novegene) is mixed with approximately 600 ng (12  $\mu$ l) of sheared genomic DNA fragments from the 2-4kb size range gel slices and 1.2  $\mu$ l of Gibco T4 ligase (5 units per  $\mu$ l) is added. Water is added to a final volume of 30 $\mu$ l and the sample is incubated overnight at 14° C.

## Transformation

The ligation reaction is titered and diluted for optimal transformation efficiency. When the ligation contains approximately 20 ng of M13 vector, the dilution will typically be from 1:25 to 1:100. A 1:25 dilution is used for plasmid ligation containing approximately 200 ng of vector DNA. To increase transformation efficiency, the ligase is denatured by heating at 65°C for 7 minutes, and placed at room temperature for 5 minutes following the heating step.

A sterile electroporation cuvette is chilled for each transformation. Electrocompetent cells are removed from the -80° C freezer and thawed on ice. For each M13 transformation, a sterile tube containing 25 ul of IPTG (25 mg/ml in water), 25  $\mu$ l of X-Gal (25 mg/ml in dimethylformamide) and 3 ml of YT top agar is prepared, capped and placed in a 45° C water bath. YT plates are pre-warmed at 37° C for several hours to avoid cross-contamination problems that may result if water remains on plates. For plasmid transformations, a sterile tube containing 0.5 ml of SOC medium is prepared for each transformation, and L + amp plates are pre-spread with 25  $\mu$ l of IPTG and 25  $\mu$ l of X-Gal.

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25 μl of electro-competent cells are mixed with DNA in diluted ligation mix in the cuvette, and the sample pulsed in an *E. coli* pulser (BioRad) set to the appropriate voltage (1.80kV for 0.1 cm cuvettes; 2.50kV for 0.2 cm cuvettes). The cuvette is removed from the pulser, and the sample immediately transferred to the tube containing SOC or YT top agar. For M13 transfections, the sample is plated immediately on YT plates. For plasmid transformations, the tube is placed in a 37° C shaker for 15-30 minutes and 30 ul aliquots are plated on L + Amp plates. Plates are incubated at 37° C overnight.

Two basic methods can be used for DNA sequencing, the chain termination method of Sanger *et al.*, *Proc. Natl. Acad. Sci. (U.S.A.)* 74:5463-5467 (1977), the entirety of which is herein incorporated by reference and the chemical degradation method of Maxam and Gilbert, *Proc. Natl. Acad. Sci. (U.S.A.)* 74:560-564 (1977), the entirety of which is herein incorporated by reference. Automation and advances in technology such as the replacement of radioisotopes with fluorescence-based sequencing have reduced the effort required to sequence DNA (Craxton, *Methods* 2:20-26 (1991), the entirety of which is herein incorporated by reference; Ju *et al.*, *Proc. Natl. Acad. Sci. (U.S.A.)* 92:4347-4351 (1995), the entirety of which is herein incorporated by reference; Tabor and Richardson, *Proc. Natl. Acad. Sci. (U.S.A.)* 92:6339-6343 (1995), the entirety of which is herein incorporated by reference). Automated sequencers are available from, for example, Pharmacia Biotech, Inc., Piscataway, New Jersey (Pharmacia ALF), LI-COR, Inc., Lincoln, Nebraska (LI-COR 4,000) and Millipore, Bedford, Massachusetts (Millipore BaseStation).

In addition, advances in capillary gel electrophoresis have also reduced the effort required to sequence DNA and such advances provide a rapid high resolution approach for sequencing DNA samples (Swerdlow and Gesteland, *Nucleic Acids Res.*. 18:1415-1419 (1990); Smith, *Nature 349*:812-813 (1991); Luckey *et al.*, *Methods Enzymol*. 218:154-172 (1993); Lu *et al.*, *J. Chromatog. A. 680*:497-501 (1994); Carson *et al.*, *Anal*.

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Chem. 65:3219-3226 (1993); Huang et al., Anal. Chem. 64:2149-2154 (1992); Kheterpal et al., Electrophoresis 17:1852-1859 (1996); Quesada and Zhang, Electrophoresis 17:1841-1851 (1996); Baba, Yakugaku Zasshi 117:265-281 (1997), all of which are herein incorporated by reference in their entirety).

A number of sequencing techniques are known in the art, including fluorescence-based sequencing methodologies. These methods have the detection, automation and instrumentation capability necessary for the analysis of large volumes of sequence data. Currently, the 377 DNA Sequencer (Perkin-Elmer Corp., Applied Biosystems Div., Foster City, CA) allows the most rapid electrophoresis and data collection. With these types of automated systems, fluorescent dye-labeled sequence reaction products are detected and data entered directly into the computer, producing a chromatogram that is subsequently viewed, stored, and analyzed using the corresponding software programs. These methods are known to those of skill in the art and have been described and reviewed (Birren *et al., Genome Analysis: Analyzing DNA*,1, Cold Spring Harbor, New York, the entirety of which is herein incorporated by reference).

PHRED is used to call the bases from the sequence trace files (<a href="http://www.mbt.washington.edu">http://www.mbt.washington.edu</a>). PHRED uses Fourier methods to examine the four base traces in the region surrounding each point in the data set in order to predict a series of evenly spaced predicted locations. That is, it determines where the peaks would be centered if there are no compressions, dropouts, or other factors shifting the peaks from their "true" locations. Next, PHRED examines each trace to find the centers of the actual, or observed peaks and the areas of these peaks relative to their neighbors. The peaks are detected independently along each of the four traces so many peaks overlap. A dynamic programming algorithm is used to match the observed peaks detected in the second step with the predicted peak locations found in the first step.

After the base calling is completed, two sequence quality steps occur 1) poor quality end sequences are cut and if the resulting sequence is 50 bp or less it is deleted 2) overall sequence quality is examined and poor sequences are deleted from the data set if

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they have an average quality cutoff below 12.5. Contaminating sequences (*E. coli*, yeast, vector, linker) are removed after sequence quality assessment.

Contigs are assembled using PANGEA clustering tools (PANGEA SYSTEMS. INC) and PHRAP (<a href="http://www.mbt.washington.edu">http://www.mbt.washington.edu</a>). PANGEA clustering tools are a series of scripts which group sequences (clusters) by comparing pairs of sequences for overlapping bases. The overlap is determined using the following high stringency parameters: word size = 8; window size = 60; and identity is 93%. Each of the clusters are then assembled using PHRAP. The final assembly output contains a collection of sequences including contigs, sequences representing the consensus sequence of overlapping clustered sequences, and singletons, sequences which are not present in any cluster of related sequences. Collectively, the contigs and singletons resulting from a DNA assembly are referred to as islands.

# Example 2

INDELs are identified by aligning sequences from Arabidopsis thaliana, Columbia and Arabidopsis thaliana, Landsberg erecta. Finished BAC sequences derived from Arabidopsis thaliana, Columbia are obtained from GenBank (http://www.ncbi.nlm.nih.gov/entrez/query.fcgi?db=Nucleotide). Because the GenBank sequences are subject to change, the finished sequences of the Arabidopsis thaliana, Columbia BACs are included herein as SEQ ID NO: 1 through SEQ ID NO: 124. The sequence of each Arabidopsis thaliana, Columbia BAC is used as a query against a database of Arabidopsis thaliana, Landsberg erecta islands using the GAP2 program of the Analysis and Annotation Tool (AAT) for Finding Genes in Genomic Sequences which was developed by Xiaoqiu Huang at Michigan Tech University and is available at the web site <a href="http://genome.cs.mtu.edu/">http://genome.cs.mtu.edu/</a>. See Huang, et al., Genomics 46: 37-45 (1997) and Huang, Computer Applications in the Biosciences 10 227-235 (1994), both of which are herein incorporated by reference in their entirety. The GAP2 program compares the query sequence with a cDNA database using a fast database search program and a rigorous alignment program. The database search program quickly identifies regions of the query sequence that are similar to a database sequence. Then the alignment program

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constructs an optimal alignment for each region and the database sequence. The output file of GAP2 is reviewed for insertions or deletions. Using alignments that are at least 96% identical (as reported by AAT), insertions and deletions are determined by looking for gaps of at least three bases, with three aligned bases on either side of the gap. To ensure that an insertion or deletion is derived from matched sequence, the 10bp region to either side of the gap is aligned and compared. To be considered an insertion or deletion, the adjacent aligned regions must be at least 90% identical (as reported by AAT). Insertions or deletions smaller than 100bp are considered candidate markers. INDELs identified by the method of this Example 2 are set forth in Table A and identified in the "method" column by reference to method 2. More particularly Table A identifies the location and nature of the polymorphism as follows:

"Seq Num" refers to the sequence of the finished BAC of *Arabidopsis thaliana*, ecotype Columbia where the polymorphism can be found;

"Seq id" refers to an arbitrary name used by applicant to identify the BAC sequence;

"Chromosome" refers to the chromosome of *Arabidopsis thaliana* in which the polymorphism is located;

"BAC Length" refers to the number of nucleotides in the finished BAC sequence; "BAC Name" refers to the name of the BAC as used in GenBank;

"Marker Name" refers to a unique six digit number arbitrarily set by applicant for a polymorphism;

"Left" refers to the position of the closest nucleotide in the flanking sequence on the 5' side of the polymorphism;

"Right" refers to the position of the closest nucleotide in the flanking sequence on the 3' side of the polymorphism;

"Type" refers to identification of the polymorphism as a SNP or IND ( $\it i.e.$ , INDEL);

"Method" refers to the method used to identify the polymorphism, where "1" represents the method of Example 3 used to detect SNPs and INDELs of less than 3

nucleotides and "2" represents the method of Example 2 used to detect large INDELs; and

"Indel Size Columbia/Landsberg" refers to the size of INDELs in terms of "n/-n" or "-n/n", where n is the size of the insertion or deletion and the minus sign indicates the ecotype with the smaller sequence length in the area of the polymorphism.

SNP Base Columbia/Landsberg" describes the nucleotide base of a SNP in the respective ecotypes, *e.g.* "T/C."

# Table A

SNP Base Columbia/ Landsberg G/A C/A G/A T/C A/T T/C A/T T/C T/A T/A
Indel Size Columbia/ Landsberg
Wethod  00000000000000000000000000000000000
Right 61034 62185 59725 62312 61731 62312 61731 62313 66146 64565 41748 42838 49676 53841 39537 59754 69645 1664 9639 16225 18886 24351 32258 35375 70795 775122 75122 75122 75122 76333 76333 76333
Left 61032 62183 59723 62310 61729 62311 66144 64563 41645 42837 49675 53840 39536 59752 69644 1662 9637 16214 18885 24350 32257 32257 32257 37251 16214 16325 76325 77325 76325 76325 76325 76325
Marker Name 468405 468406 468406 468409 468409 468409 468410 468575 468575 470714 470715 470717 471481 470715 469902 469903 469909 469909 469910 469911 469910 469911 469911 469911 469911 469911 469911 469911 469911 469911
BAC Name TIN6 TIN6 TIN6 TIN6 TIN6 TIN6 TIN6 TIN6
BAC Chromosome Length 76175 1 76175 1 107234 1 107234
Seq id AC009273 AC007583
Seg

SNP Base Columbia/ Landsberg	C/A C/A A/G	CT CT CT G/A G/C	I/G G/C A/G A/T C/T	77.4 A G G/A C/T
Indel Size Columbia/ Landsberg 5/-5 -4/4	CIC-			-6/6 -3/3 -54/54
Method 2	7			
Type IND IND	SNP SNP SNP SNP	SNP SNP SNP SNP SNP	SNP SNP SNP SNP	S S S S S S S S S S S S S S S S S S S
Right 86491 96587	26802 26919 27393	26753 26753 48519 48651 12023 12525	45189 65812 65767 67898 68223 67897	24352 64002 42908 14868 15602 15554 729 729 729 729 47145 46889 46889 46889 46889 21545 21545 2826 30481
Left 86485 96586	26800 26917 27391	26751 26751 48517 48649 12021 12523	45187 65810 65765 67896 68221 67895	24350 64000 42906 14866 15600 15552 727 793 108 47143 46886 47021 46886 47021 46889 21544 28358 30476
Marker Name 469919 469920	469921 472002 472003 472004	472005 472241 472242 472372 472373	472687 472745 472746 472936 472937	473013 473013 473037 473433 473434 473579 473520 473620 473622 473622 473622 473622 473622 473623 473623 473623 473624 473623
BAC Name F24B9 F24B9	F24B9 F13K23 F13K23 F13K23	F13K23 F13K23 F13K23 F13K23 F13K23	F13K23 F13K23 F13K23 F13K23 F13K23	F13K23 F13K23 F13K23 F13K23 F13K23 F13K23 F13K23 F13K23 F13K23 F13K23 F13K23 F13K23 F13K23 F13K23 F13K23 F13K23
BAC Chromosome Length 1 107234	1 10/234 1 74328 1 74328 1 74328	1 74328 1 74328 1 74328 1 74328 1 74328	1 74328 1 74328 1 74328 1 74328 1 74328	1 74328 1 74328
Seq id AC007583 AC007583	AC00/583 AC012187 AC012187 AC012187	AC012187 AC012187 AC012187 AC012187 AC012187	AC012187 AC012187 AC012187 AC012187 AC012187	AC012187
Seq num 3	w 4 4 4	1 4 4 4 4 4	4 4 4 4 4 4	1 4 4 4 4 4 4 4 4 4 4 4 4 4 4 4 4 4 4 4

SNP Base Columbia/	Landsberg										A/C	T/C	G/T	A/T	A/T	T/C	. Q/A	A/T	A/T	T/G	A/G	C/A	T/C	A/G	T/A	G/A	T/A	T/C	T/C	A/C	T/C	A/G	G/T	A/T	G/T	C/T	T/C
Indel Size Columbia/	Landsberg	3/-3	12/-12	-3/3	-2/2	-1/1	-2/2	-1/1	3/-3	-1/1																											
	Method	2	2	2	_	1	1	-	=	1	1	1	I	<b>—</b>		-	-	-		1	1	1	1	ı	-	1	_	1	П		1	-	1	1	1	1	1
	Type	QNI	IND	ONI	IND	IND	ONI	ONI	ONI	ONI	SNP																										
	Right	31807	31872	350	14242	24021	355	43083	46896	73894	8013	8142	8012	38077	39804	5899	68784	84818	85087	50659	80331	12167	12014	12591	16374	16763	17422	14587	15164	16366	13254	13347	14316	15640	13215	16186	56853
	Left	31803	31859	349	14241	24020	354	43082	46892	73893	8011	8140	8010	38075	39802	5897	68782	84816	85085	50657	80329	12165	12012	12589	16372	16761	17420	14585	15162	16364	13252	13345	14314	15638	13213	16184	56851
Marker	Name	473825	473826	473827	474047	474048	474049	474050	474051	474052	467114	467115	467116	467979	467980	468122	468123	468637	468638	468662	468726	468752	468753	468754	468755	468756	468757	468758	468759	468760	468761	468762	468763	468764	468765	468766	468883
BAC	Name	F13K23	F14L17																																		
BAC	Chromosome Length	74328	74328	74328	74328	74328	74328	74328	74328	74328	111686	111686	111686	111686	111686	111686	111686	111686	111686	111686	111686	111686	111686	111686	111686	111686	111686	111686	111686	111686	111686	111686	111686	111686	111686	111686	111686
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	Seq id	AC012187	AC012188																																		
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SNP Base Columbia/ Landsberg G/C A/G A/T T/C C/G C/T C/G C/T A/G A/C A/T T/G G/A G/A G/A A/T	
Indel Size Columbia/ Landsberg 3/-3 3/-3 3/-3 3/-3 3/-3 13/-13 4/-4 -8/8 18/-18 -2/2 1/-1 -1/1 -1/1 -1/1	-62/62 4/-4 -3/3 9/-9
Method	0000
SNP	
Right 10115 44313 44329 28004 28266 28436 12921 12998 25376 53939 550 7169 13242 13242 13242 13242 10031 17977 1758 17955 34824 63540 63635 69438	1645 43324 4664 53393
Left 10113 44311 44321 28002 28264 28434 12917 12994 25080 25362 53934 589 13218 13229 13241 13259 28090 52197 8237 28529 19140 10029 57756 17975 17953 69633	1644 43319 4663 53383
Marker Name 468970 469164 469165 469228 469229 469229 469641 469644 469644 469645 469645 469647 471133 471133 471133 471133 471133 467267 467267 467283 468269 468398	470620 470621 470622 470623
BAC Name F14L17 F16N11 T16N11 T16N11 T16N11 T16N11 T16N11 T16N11 T16N11	T16N11 T16N11 T16N11 T16N11
BAC Chromosome Length 111686	91001 91001 91001 91001
	453 1 453 1 453 1 453 1
Seq id     AC012188     AC012453     AC013453     AC013453     AC013453     AC013453	AC013453 AC013453 AC013453 AC013453
Seq num	9 9 9

Columbia/ Landsberg Ly/A A/G C/T T/A G/A G/A G/A G/A G/A C/A C/A C/A C/T	
Columbia/ Landsberg 4/-4 -9/9 4/-4 -3/3 -3/3 -3/3 -40/40 38/-38 -10/10 -4/4 -7/7 -4/4 1/-1 1/-1 1/-1	ı i
Method	•
	!
Right 53812 54080 61857 65846 74185 75367 824 82377 87653 89602 89502 89518 17209 57920 57920 57920 57920 57920 57920 57920 5793 81752 20429 20429 20429 30594 30594	) ) 
Left 53807 54079 54245 61856 65845 73501 75366 785 82376 83456 89601 89517 17207 57919 58215 38649 38649 38649 38649 38649 38649 20427 2791 2665 1050 30692	: : :
Marker Name 470624 470625 470625 470626 470629 470629 470631 470631 470631 470633 470633 470633 470633 470633 470633 470634 470633 471972 471972 471972 471972 471972 471972 471973 473514 473514 473575 473576 473576 473576	) - -
BAC Name T16N111 T16N11 T16	
BAC Chromosome Length 1 1 91001 1 91001 1 91001 1 91001 1 91001 1 91001 1 91001 1 91001 1 91001 1 84974 1 84974	/10
Seq id AC013453 AC007843	710000
Seq num e e e e e e e e e e e e e e e e e e e	-

SNP Base Columbia/ Landsberg	1/C G/A A/G 1/C G/A C/G	C/G A/G C/A A/G C/T C/T C/A C/A C/A C/A C/A C/A C/A C/A C/A
Indel Size Columbia/ Landsberg -1/1 -3/3 -1/1 1/-1 -1/1 -5/5		3/-3
Method 1 1 1 1 1 1 1 1 1		
Type IND UNI UNI UNI UNI UNI UNI UNI	SNP SNP SNP SNP SNP SNP	SNP
Right 32466 38916 50806 63016 68447 74023	10921 11649 11668 88015 28403 28012 28435	28435 75707 50260 50361 49637 86165 51422 13893 13688 61183 60466 24704 59365 17438 80514 80514 80287 80287 80204 80287 80401 89145 89145
Left 32465 38915 50805 63014 68446 74022 82176	10919 11647 11666 88013 28401 28010	28433 75705 50258 50359 49635 86163 51420 13891 13886 61181 60464 24702 59363 17436 80512 80469 80302 80302 80309 80143 89143
Marker Name 474169 474171 474171 474172 474173 474173	466842 466843 466844 467047 467102 467103	467104 467137 467173 467173 467174 467329 467329 467537 467950 468999 468999 468527 468527 468529 468530 468530 468530 468530 468530 468530 468530 468836
BAC Name F28G4 F28G4 F28G4 F28G4 F28G4 F28G4	F15H18 F15H18 F15H18 F15H18 F15H18 F15H18	F15H18
BAC Chromosome Length	95327 95327 95327 95327 95327 95327 95327	95327 95327 95327 95327 95327 95327 95327 95327 95327 95327 95327 95327 95327 95327 95327
Seq id CO07843 1 AC007843 1 AC007843 1 AC007843 1 AC007843 1 AC007843 1 AC007843 1	AC013354 1 AC013354 1 AC013354 1 AC013354 1 AC013354 1 AC013354 1	AC013354
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Indel Size

Columbia/	Landsberg													G/A	A/G	I/G	I/G	I/G	G/T	C/G	I/C	C/T	I/C	I/A	G/A	G/T	A/G	C/A	A/G	A/G	A/T	G/A	G/A	T/G	T/A	G/A
_	Landsberg I	-13/13	-5/5	-5/5	-5/5	-3/3	9 <i>L</i> /9 <i>L</i> -	-3/3	9-/9	-1/1	-1/1	-2/2			7				•			•				•										
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	Right	34120	34719	34729	34730	799	69756	70000	93740	49808	51699	60964	62793	78436	1727	99380	99590	99632	99631	10627	90920	91263	93643	11983	61145	71036	100767	41774	41840	41794	41792	48683	48037	48013	12697	12712
	Left	34119	34718	34728	34729	661	69755	66669	93733	49807	51698	69609	62791	78434	1725	99378	88566	99630	99629	10625	90918	91261	93641	11981	61143	71034	100765	41772	41838	41792	41790	48681	48035	48011	12695	12710
Marker	Name	469687	469688	469689	469690	469691	469692	469693	469694	471151	471152	471153	471154	466830	467071	467516	467517	467518	467519	467636	467909	467910	467912	467956	467996	468404	468730	468930	468931	468932	468933	469327	469328	469329	469359	469360
BAC	Name	F15H18	F15H18	F15H18	F15H18	F15H18	F15H18	F15H18	F15H18	F15H18	F15H18	F15H18	F15H18	F6A14																						
BAC	Chromosome Length		95327	95327	95327	95327	95327	95327	95327	95327	95327	95327	95327	108767	108767	108767	108767	108767	108767	108767	108767	108767	108767	108767	108767	108767	108767	108767	108767	108767	108767	108767	108767	108767	108767	108767
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	Seq id	AC013354	AC013354	AC013354	AC013354	AC013354	AC013354	AC013354	AC013354	AC013354	AC013354	AC013354	AC013354	AC011809																						
Sec	_		∞	∞	<b>∞</b>	∞	∞	∞	∞	∞	∞	∞	∞	6	6	6	6	6	6	6	6	6	6	6	6	6	6	6	6	6	6	6	6	6	6	6

SNP Base Columbia/ Landsberg	A/T A/T																										-								C/G
Indel Size Columbia/ Landsberg		4/-4	5/-5	8-/8	5/-5	5/-5	-4/4	-5/5	L-/L	3/-3	26/-26	14/-14	14/-14	-21/21	L/L-	3/-3	3/-3	47/-47	-4/4	8-/8	14/-14	16/-16	5/-5	8/8-	16/-16	3/-3	-5/5	-2/2	-1/1	3/-3	1/-1	-1/1	-1/1	1/-1	
Method	<b>-</b>	2	2	2	7	2	2	2	2	2	2	2	7	2	2	7	2	2	2	2	2	2	2	2	2	2	2	<del></del>			1	1	1	1	1
Type	SNP SNP	ONI	ON N	ONI	ONI	ONI	IND	ONI	IND	ONI	ONI	ONI	IND	ONI	ONI	ONI	ONI	ONI	ONI	IND	ON N	ONI	ONI	SNP											
Right	12752 12885	1233	17420	17548	19487	19490	26342	29070	3634	41746	49772	51391	52587	67268	69172	60309	69413	72592	72702	79638	79735	80807	80935	81548	82702	83461	84373	26347	35006	41752	41843	70953	71085	89200	33215
Left	12750 12883	1228	17414	17539	19481	19484	26341	53063	3626	41742	49745	51376	52572	67267	69171	69305	69409	72544	72701	79629	79720	80790	80929	81547	82685	83457	84372	26346	35005	41748	41841	70952	71084	89198	33213
Marker Name	469362 469363	470349	470350	470351	470352	470353	470354	470355	470356	470357	470358	470359	470360	470361	470362	470363	470364	470365	470366	470367	470368	470369	470370	470371	470372	470373	470374	471351	471352	471353	471354	471355	471356	471357	472001
BAC Name	F6A14 F6A14	F6A14	F6A14	F6A14	F6A14	F6A14	F6A14	F6A14	F6F9																										
BAC Chromosome Length	108767 108767	108767	108767	108767	108767	108767	108767	108767	108767	108767	108767	108767	108767	108767	108767	108767	108767	108767	108767	108767	108767	108767	108767	108767	108767	108767	108767	108767	108767	108767	108767	108767	108767	108767	119942
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Seq id	AC011809 AC011809	AC011809	AC011809	AC011809	AC011809	AC011809	AC011809	AC011809	AC007797																										
Seq	6 6	. 6	6	6	6	6	6	6	6	6	6	6	6	6	6	6	6	6	6	6	6	6	6	6	6	6	6	6	6	6	6	6	6	6	10

Indel Size

Columbia/	Landsberg	$C/\Gamma$	G/A	A/G	CT	A/C	A/C	T/G	A/T	G/T	T/C	A/T	A/T	T/A	G/T	T/C	C/A	C/A	G/A	T/A	A/C	T/G	A/G	CT	G/A	C/A	CT	T/A	G/A	T/C	G/C	A/T	T/A	CT			
Columbia/	Landsberg																																		-3/3	6-/6	9-/9
	Method	1	1	1	_	<b>,</b>	1	1	_	-		_	-	1	_	1			_	1	1	1		_		1	1	-	-		_		1	-	2	2	_
	Type	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	ONI		ONI
	Right	31296	109306	108721	13285	28436	28253	28570	52874	59392	39441	45351	45350	3654	19500	35108	50552	50616	50721	50801	50551	50682	50958	50527	76751	76406	76468	71612	22855	23988	26842	26841	22293	21994	1379	1646	24169
	Left	31294	109304	108719	13283	28434	28251	28568	52872	59390	39439	45349	45348	3652	19498	35106	50550	50614	50719	50799	50549	20680	50956	50525	76749	76404	76466	71610	22853	23986	26840	26839	22291	21992	1378	1636	24162
Marker	Name	472026	472281	472282	472340	472341	472342	472343	472405	472418	472419	472764	472765	472790	472791	472797	472804	472805	472806	472807	472808	472809	472810	472811	473122	473123	473124	473162	473279	473280	473355	473356	473707	473708	473923	473924	474273
BAC	Name	F6F9	F6F9	F6F9	F6F9	F6F9	F6F9	F6F9	F6F9	F6F9	F6F9	F6F9	F6F9	F6F9	F6F9	F6F9	F6F9	F6F9	F6F9	F6F9	F6F9	F6F9	F6F9	F6F9	F6F9	F6F9	F6F9	F6F9	F6F9	F6F9	F6F9	F6F9	F6F9	F6F9	F6F9	F6F9	F6F9
BAC	Chromosome Length		119942	119942	119942	119942	119942	119942	119942	119942	119942	119942	119942	119942	119942	119942	119942	119942	119942	119942	119942	119942	119942	119942	119942	119942	119942	119942	119942	119942	119942	119942	119942	119942	119942	119942	119942
	Chro	1	_	_	_	1	_		1	1	Т	1	1	1	1	-	_	1	1	1	_	1	-	1	_	_	_	_	_	_	_	_		_	_	_	1
	Seq id	AC007797	AC007797	AC007797	AC007797	AC007797	AC007797	AC007797	AC007797	AC007797	AC007797	AC007797	AC007797	AC007797	AC007797	AC007797	AC007797	AC007797	AC007797	AC007797	AC007797	AC007797	AC007797	AC007797	AC007797	AC007797	AC007797	AC007797	AC007797	AC007797	AC007797	AC007797	AC007797	AC007797	AC007797	AC007797	AC007797
S.	T III	10	10	10	10	10	10	10	10	10	10	10	10	10	10	10	10	10	10	10	10	10	10	10	10	10	10	10	10	10	10	10	10	10	10	10	10

SNP Base Columbia/ Landsberg	G/C C/G G/A 17/C 17/A G/A	G/A G/A G/A G/T A/T A/C	AT G/A C/A A/G G/A
Indel Size Columbia/ Landsberg -1/1 1/-1		-5/5 -5/5 3102/-3102 3101/-3101 -3/3 -6/6 14/-14 29/-29 -1/1	
Method 1 1 1	·		
Type IND IND IND	SNP SNP SNP SNP SNP SNP SNP SNP		SNP SNP SNP SNP
Right 45597 50572 50802	45549 45147 4238 4239 4058 36572 31731	31681 31686 31577 31716 313289 33302 16067 16068 28287 28349 33225 34493 34493 34609 41402 62144 62145	41665 43454 21557 21631 54383
Left 45596 50570 50800 71576	45547 45145 4236 4237 4056 36570 31729	31679 31679 31575 31714 31680 33287 33300 16066 16067 25247 33224 34492 34492 34608 41387 62114 62115 3797	41663 43452 21555 21629 54381
Marker Name 474274 474275 474276	466902 466903 467340 467341 468248 469064	469066 469067 469068 469069 469080 469181 470825 470825 470826 470829 470833 470833 470833 470833 470833	471693 471937 472249 472250 472427
BAC Name F6F9 F6F9 F6F9	72211 722111 722111 722111 722111 722111	72211 722111 722111 722111 722111 722111 722111 722111 722111 722111 722111 722111	F2418 F2418 F2418 F2418 F2418
BAC Chromosome Length 1 119942 1 119942 1 119942	98689 98689 98689 98689 98689 98689 98689	98689 98689 98689 98689 98689 98689 98689 98689 98689 98689 98689 98689 98689	108365 108365 108365 108365
Chr 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1			
Seq id AC007797 AC007797 AC007797	AC012190 AC012190 AC012190 AC012190 AC012190 AC012190	ACO12130 ACO12190 ACO12190 ACO12190 ACO12190 ACO12190 ACO12190 ACO12190 ACO12190 ACO12190 ACO12190 ACO12190 ACO12190 ACO12190	AC015447 AC015447 AC015447 AC015447 AC015447
Seq num 10 10 10	2=======	=======================================	12 12 12 12 12 12 12 12 12 12 12 12 12 1

SNP Base Columbia/	Landsberg	1/C	Q	J/L	C/L	G/C	G/A	D/O	G/A	T/C	A/C	A/G	A/G	T/G	A/G	A/G	A/G	A/G	C/T	T/A	A/G	C/A	C/A	G/A	T/A	A/G									T/A	T/C
Indel Size Columbia/	Landsberg																										-5/5	-3/3	6-/6	1/-1	1/-1	1/-1	1/-1	1/-1		
,	Method	<b>→</b>	<b>-</b> -	<b>-</b> -	- <del>-</del>	·	_	1	1	1	_	_	П	_	1	1	1	_	-	_	_			-		П	2	2	2	-	_	_			-	-
	Type	SNF	SNF	CIVIS	SNP TNP	SNP	ON.	ON N	QNI	ONI	ONI	ONI	ONI	ONI	SNP	SNP																				
	Right	34338 0508	930o 430	5,50	3/2 1163	7859	97727	81703	39368	39823	41086	39868	40295	40914	41344	39254	39365	39409	39906	55350	55868	18973	11022	16318	88377	88139	1714	2951	3809	21782	21792	54580	54582	88479	80056	38143
	Left	34330 0506	9300	975	1161	7857	97725	81701	39366	39821	41084	39866	40293	40912	41342	39252	39363	39407	39904	55348	55866	18971	11020	16316	88375	88137	1713	2950	3799	21780	21790	54578	54580	88477	80054	38141
Marker	Name	472428	472046	770040	472947	472971	473018	473069	473074	473075	473076	473077	473078	473079	473080	473081	473082	473083	473084	473333	473334	473359	473448	473706	473800	473801	473876	473877	473878	474151	474152	474153	474154	474155	466960	467546
BAC	Name	F24J8	F24J8	F2410	F24J8 F24J8	F2418	F24J8	F2418	F24J8	F24J8	F24J8	F24J8	F24J8	F24J8	T26F17	T26F17																				
BAC		108365	108365	106303	108365	108365	108365	108365	108365	108365	108365	108365	108365	108365	108365	108365	108365	108365	108365	108365	108365	108365	108365	108365	108365	108365	108365	108365	108365	108365	108365	108365	108365	108365	82875	82875
	Chromosome	<b>-</b> -	<b>-</b> -	<b>⊣</b> -		<b>-</b>	·	-	_	_	1	_	Ţ	_	_			_	1	_		_	_	-	1	T	_	1	<del></del>		_	_	_	1	-	-
	Seq id	AC015447	AC015447	AC015447	AC01544/ AC015447	AC015447	AC015447	AC015447	AC015447	AC015447	AC015447	AC013482	AC013482																							
Seq	mnu	2 5	2 :	71	7 2	2 2	12	12	12	12	12	12	12	12	12	12	12	12	12	12	12	12	12	12	12	12	12	12	12	12	12	12	12	12	13	13

SNP Base Columbia/	Landsberg	T/C	C/T	C/A	T/C	T/C	G/A	T/C	C/A	C/T	A/G	T/A	T/A	G/C	A/G	AT	A/T	CT	A/C	G/T	C/T	A/T	T/C														
Indel Size Columbia/	Landsberg																							-5/5	-4/4	-3/3	-4/4	3/-3	3/-3	236/-236	236/-236	546/-546	9/9-	8/8-	-3/3	36/-36	4/-4
	Method			-	_	1		_	1	<del></del> 1	_	1	_	_		_	_	-		1	-		_	2	2	2	2	2	2	2	2	2	2	2	2	2	2
	Type	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SINP	ONI	ONI	ONI	ONI	IND	ONI	ONI	ONI	ONI	ONI	ONI	ONI	ONI	ONI								
	Right	38131	38273	68895	68918	39176	50415	50078	17850	17995	13243	14583	14565	16648	15926	14622	14570	16099	5140	5084	5085	48139	52138	11724	15918	15942	36133	39862	48269	65495	65497	71656	72095	72391	7477	74964	77559
	Left	38129	38271	68893	68916	39174	50413	50076	17848	17993	13241	14581	14563	16646	15924	14620	14568	16097	5138	5082	5083	48137	52136	11723	15917	15941	36132	39858	48265	65258	65260	71109	72094	72390	7476	74927	77554
Marker	Name	467547	467548	467599	467600	467873	467937	467938	468489	468490	468491	468492	468493	468494	468495	468496	468497	468498	468533	468534	468535	468554	469195	470857	470858	470859	470860	470861	470862	470863	470864	470865	470866	470867	470868	470869	470870
BAC	Name	T26F17	T26F17	T26F17	T26F17	T26F17	T26F17	T26F17	T26F17	T26F17	T26F17	T26F17	T26F17	T26F17	T26F17	T26F17	T26F17	T26F17	T26F17	T26F17	T26F17	T26F17	T26F17	T26F17	T26F17	T26F17	T26F17	T26F17	T26F17								
BAC	Chromosome Length	82875	82875	82875	82875	82875	82875	82875	82875	82875	82875	82875	82875	82875	82875	82875	82875	82875	82875	82875	82875	82875	82875	82875	1 82875	1 82875	1 82875	1 82875	1 82875	1 82875	1 82875	1 82875	1 82875	1 82875	1 82875	1 82875	1 82875
	Seq id C	AC013482 1	AC013482 1	AC013482 1	AC013482 1	AC013482 1	AC013482 1	AC013482 1	AC013482 1	AC013482 1	AC013482 1	AC013482	AC013482 1	AC013482	AC013482 1	AC013482 1	AC013482																				
Sed	_	13	13	13	13	13	13	13	13	13	13	13	13	13	13	13	13	13	13	13	13	13	13	13	13	13	13	13	13	13	13	13	13	13	13	13	13

SNP Base Columbia/ Landsberg	G/A C/A T/C A/G C/T T/C C/T T/C G/A G/A
Indel Size Columbia/ Landsberg 4/-4 -3/3 -8/8 -1/1 1/-1 -4/4 -3/3 1/-1 2/-2 1/-1 2/-2 1/-1 -4/4	-3/3
Method	
Type CN CN C	S S S S S S S S S S S S S S S S S S S
Right 77588 78802 78963 14573 14573 14584 15921 15950 26080 39364 39867 48268 48270 68896	54816 66532 66054 65685 66535 55302 67254 79952 34008 20670 19755 19125 68134 83187 81292 37753 21863
Left 77583 78801 78962 14572 14582 15920 15920 15949 26078 39362 39861 39865 48265 48268 68895	54814 66530 66533 66533 66533 55500 55321 67252 79950 34006 20668 19753 19123 68176 68132 83185 81290 37751 21862
Marker Name 470871 470872 470873 471543 471544 471545 471546 471549 471551 471553 471553	471730 471756 471757 471758 471759 471936 472331 472759 472759 472760 472761 472760
BAC Name T26F17	T1K7 T1K7 T1K7 T1K7 T1K7 T1K7 T1K7 T1K7
BAC Chromosome Length 82875 1 82875 1 82875	89473 89473 89473 89473 89473 89473 89473 89473 89473 89473 89473 89473 89473 89473 89473 89473
	AC013427 1
B B	4     4

SNP Base Columbia/ Landsberg	1/C 1/G 1/G C/T C/T	G/A A/T C/G C/G G/A A/G A/G C/T C/T C/G
Indel Size Columbia/ Landsberg 1/-1 1/-1 1/-1 -1/1 -5/5	·	822/-822 -10/10 -9/9 -9/9 11/-11 3/-3
Method 1 1 1 1 1		
Type GNI GNI GNI GNI CNI CNI CNI CNI CNI CNI CNI CNI CNI C	S S S S S S S S S S S S S S S S S S S	S S S S S S S S S S S S S S S S S S S
Right 29409 29411 38825 5749 5751	42169 44295 44360 44078 78197 49206 50899	50899 39954 40342 38307 74113 79657 70427 71353 70757 51954 76717 76883 105249 105143 102193 102997 102999 3030 34265
Left 29407 29409 38823 5748 5750	44293 44293 44358 44076 44091 78195 49204	50897 39952 40340 38305 74111 79655 70425 71351 70755 51952 76715 76881 105247 105247 102330 102996 102998 3018 34261
Marker Name 474408 474409 474410 474411	466845 466846 466847 466849 467088 467194	467195 467724 467725 467725 467757 468870 468871 469192 469193 469193 469194 469194 469197 469198 469246 469246 469247 470767 470770 470770
BAC Name T1K7 T1K7 T1K7 T1K7	122C5 122C5 122C5 122C5 122C5 122C5 122C5	122C5 122C5
BAC Chromosome Length 1 89473 1 89473 1 89473 1 89473 1 89473	1 106753 1 106753 1 106753 1 106753 1 106753 1 106753 1 106753	1 106753 1 106753
Seq id AC013427 AC013427 AC013427 AC013427	AC012375 AC012375 AC012375 AC012375 AC012375 AC012375	AC012375 AC012375 AC012375 AC012375 AC012375 AC012375 AC012375 AC012375 AC012375 AC012375 AC012375 AC012375 AC012375 AC012375 AC012375 AC012375 AC012375 AC012375 AC012375
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Columbia/	Landsberg	-4/4	-4/4	-3/3	L-1L	4/-4	2/-2	3/-3	-5/5	32/-32	64/-64	64/-64	6/6-	6/6-	8/8-	6/6-	6/6-	1271/-12	5/-2	905/-905	-3/3	-1/1	-2/2	1/-1	2/-2	2/-2	1/-1										
	Method	2	2	2	2	2	2	2	2	2	2	2	2	2	2	2	2	2	2	2		_	_	<b>—</b>	1	_	1	1				<b>-</b>	1		-	1	_
	Type	ONI	ONI	IND	ONI	ONI	ONI	IND	ONI	ONI	IND	ONI	ONI	IND	ONI	ONI	ONI	ONI	IND	ONI	IND	ON.	ONI	ONI	ONI	QNI	ONI	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP
	Right	3766	3767	40489	46979	48108	73994	88705	88745	88912	88944	88978	92764	92769	93018	93019	93020	94398	009/6	69666	40491	52423	73914	73990	73993	73996	80254	78932	79397	79011	78931	79148	75458	7910	7923	8543	10549
	Left	3765	3766	40488	46971	48103	73988	88701	88744	88879	88879	88913	92763	92768	93017	93018	93019	93126	97594	99063	40490	52422	73913	73988	73990	73993	80252	78930	79395	79009	78929	79146	75456	7908	7921	8541	10547
Marker	Name	470774	470775	470776	470777	470778	470779	470780	470781	470782	470783	470784	470785	470786	470787	470788	470789	470790	470791	470792	471511	471512	471513	471514	471515	471516	471517	466986	466987	466988	466989	466990	467048	467089	467090	467091	467092
BAC	Name	T22C5	T22C5	T22C5	T22C5	T22C5	T22C5	T22C5	T22C5	T22C5	T22C5	T22C5	T22C5	T22C5	F3M18	F3M18	F3M18	F3M18	F3M18	F3M18	F3M18	F3M18	F3M18	F3M18													
BAC	Chromosome Length	106753	106753	106753	106753	106753	106753	106753	106753	106753	106753	106753	106753	106753	106753	106753	106753	106753	106753	106753	106753	106753	106753	106753	106753	106753	106753	104163	104163	104163	104163	104163	104163	104163	104163	104163	104163
	Chrom	1	_		_		_				_	1	_	_	<del></del>		_	~	_	_	_	_	_	_	1	1	_	-	1	1	_		1	1	_	1	_
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Sed	unu	15	15	15	15	15	15	15	15	15	15	15	15	15	15	15	15	15	15	15	15	15	15	15	15	15	15	16	16	16	16	16	16	16	16	16	16

SNP Base Columbia/	Landsberg	C/T	T/A	A/G	A/C	C/T	C/T	A/T	T/A	T/C	G/C	C/G	A/T	G/A	A/C	T/C	A/T	G/T	A/G	C/T	A/C	T/C	C/T	C/T	G/C	T/G	G/C	T/C	G/A	G/A	T/A	G/C	T/C	T/G	C/G	A/T	G/A
Indel Size Columbia/	Landsberg																																				
	Method	1	-		-	-	1	1	1	_	_	1	1	1	1	1			1	-1	_		-	1	_	1		1	1	1	-	_	_	_	<b>—</b>	_	
	Type	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP																
	Right	10697	76531	77227	60553	60291	8/009	61428	62434	63353	63199	63331	62377	56890	57077	57377	58015	57317	44900	44962	97539	68696	98406	98405	90837	91323	55309	56176	66069	66195	65630	66772	66215	66579	67032	65836	94806
	Left	10695	76529	77225	60551	60289	92009	61426	62432	63351	63197	63329	62375	56888	57075	57375	58013	57315	44898	44960	97537	28696	98404	98403	90835	91321	55307	56174	26069	66193	65628	0//99	66213	66577	67030	65834	94804
Marker	Name	467093	467225	467226	467477	467478	467479	467480	467481	467482	467483	467484	467485	467841	467842	467843	467844	467845	467904	467905	468012	468013	468101	468102	468389	468390	468880	468881	469149	469150	469151	469152	469153	469154	469155	469156	469268
BAC	Name	F3M18	F3M18	F3M18	F3M18	F3M18	F3M18	F3M18	F3M18	F3M18	F3M18	F3M18	F3M18	F3M18	F3M18	F3M18	F3M18	F3M18	F3M18	F3M18	F3M18																
BAC	Chromosome Length	104163	104163	104163	104163	104163	104163	104163	104163	104163	104163	104163	104163	104163	104163	104163	104163	104163	104163	104163	104163	104163	104163	104163	104163	104163	104163	104163	104163	104163	104163	104163	104163	104163	104163	104163	104163
	Chro	_			_	_	П	_	_	_	_	_	_	_	1			<del></del>	_	_	Т	_	_	_	_	<del></del>		_	_	_	_	_		_	_	_	П
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Sea	unu	16	16	16	16	16	16	16	16	16	16	16	16	16	16	16	16	16	16	16	16	16	91	16	16	16	16	16	16	16	91	16	16	16	16	16	16

SNP Base Columbia/ Landsberg T/C T/C	A/C C/G C/T C/A A/T A/T A/T C/G C/G C/G C/T T/A C/T C/G C/T C/T C/G C/T C/G C/T C/G C/T C/G C/T C/G C/T C/T C/T C/T C/T C/T C/T C/T C/T C/T
Indel Size Columbia/ Landsberg -3/3 2664/-2664 3/-3 4/-4 10/-10 -11/11 -3/3 6/-6 -23/23 -6/6 7/-7 1/-1 1/-1	
Method 1 2 2 2 2 1 1 1 1 1	
Type SNP SNP CN	SAP SAP SAP SAP SAP SAP SAP SAP SAP SAP
Right 95462 94869 15914 23958 29837 45686 46220 53859 62646 68688 72675 87257 87832 31103 55220 69183	61405 61405 61460 61836 61808 113790 114406 89243 89441 115706 115881 81716 124362 12455 124655 124655 96429
Left 95460 94867 15913 21293 29833 45681 46209 53858 62645 68681 72674 87256 87256 87256 87256	69194 61403 61458 61834 61806 113788 114404 89441 89441 89279 64439 115704 115879 81714 124360 124360 12453 96427 96427
Marker Name 469269 469270 470267 470268 470271 470272 470273 470277 471339 471339	471748 471749 471750 471751 471785 471785 471885 471885 471885 471856 471957 472028 472027 472028 472027 472028 472027 472028
BAC Name F3M18	F3M18 F28N24
BAC Chromosome Length 1 104163 1 104163 1 104163 1 104163 1 104163 1 104163 1 104163 1 104163 1 104163 1 104163 1 104163 1 104163 1 104163	1 104.103 1 154716 1 154716
Seq id AC010155 AC010155 AC010155 AC010155 AC010155 AC010155 AC010155 AC010155 AC010155 AC010155 AC010155 AC010155 AC010155	AC010155 AC021043
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SNP Base Columbia/ Landsberg T/C A/G A/G G/A G/A T/A G/C	A/G A/G T/A C/T G/A G/A C/G
Indel Size Columbia/ Landsberg -1/1 1/-1 1/-1 1/-1 1/-1 1/-1 -1/1 -1/	1,-1 -1/1 1,-1 1/-1
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Right 48847 48911 58533 49474 49610 83521 100180 10383 13372 19561 38587 49405 69494 89353 89354 89353 89354 89353	96496 97505 97589 99883 28154 28228 46454 2250 13358 13421 13380
Left 48845 48909 58531 49472 49608 83519 100157 100305 100178 10375 10381 13369 19559 38585 49404 69487 89353 89405 90927 90620	96494 97504 97587 99881 28152 28226 46452 2248 13356 13378 13183
Marker Name 472542 472543 473127 473192 473193 473557 473559 474184 474185 474189 474190 474190 474191 474192 474192 474192 474193 474194 474194	474198 474199 474200 474201 471676 471677 471830 471830 471909 471909
BAC Name F28N24	F28N24 F28N24 F28N24 F28N24 F5D14 F5D14 F5D14 F5D14 F5D14 F5D14
BAC  bac  length  ls4716  ls4716	154716 154716 154716 154716 127462 127462 127462 127462 127462 127462
Chromosome 1	
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SNP Base Columbia/	Landsberg	A/T	G/T	G/A	G/C	T/C	C/T	C/T	G/A	T/C	A/T	CT	G/T	T/A	T/A	G/A	T/A	T/A	T/C	T/C	A/G	A/T	A/T	C/T	I/C	C/T	G/T	T/C	C/A	A/G	G/A	C/A	C/A	G/A	T/A	G/A	T/A
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	Right	13177	13379	65928	65672	65617	24935	24919	9360	64444	64142	126440	41745	37147	34840	34842	34850	34852	36707	34841	34846	37188	34844	34847	39498	76303	76595	95188	16397	16961	21709	78423	79601	79599	79490	79128	79124
	Left	13175	13377	65926	02959	65615	24933	24917	63958	64442	64140	126438	41743	37145	34838	34840	34848	34850	36705	34839	34844	37186	34842	34845	39496	76301	76593	95186	16395	16959	21707	78421	79599	79597	79488	79126	79122
Marker	Name	471912	471913	471949	471950	471951	472121	472122	472314	472315	472316	472389	472988	473051	473052	473053	473054	473055	473056	473057	473058	473059	473060	473061	473121	473487	473488	473528	473577	473578	473702	473723	473724	473725	473726	473727	473728
BAC	Name	F5D14	F5D14	F5D14	F5D14	F5D14	F5D14	F5D14	F5D14	F5D14	F5D14	F5D14	F5D14	F5D14	F5D14	F5D14																					
BAC	Chromosome Length	127462	127462	127462	127462	127462	127462	127462	127462	127462	127462	127462	127462	127462	127462	127462	127462	127462	127462	127462	127462	127462	127462	127462	127462	127462	127462	127462	127462	127462	127462	127462	127462	127462	127462	127462	127462
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SNP Base Columbia/ Landsberg	1/C G/C	T/G														A/T	G/A	T/A	C/A	C/G															
Indel Size Columbia/ Landsberg			-3/3	-4/4	11-7	-4/4	-2/2	-3/3	-1/1	-3/3	2/-2	1/-1	2/-2	1/-1	-1/1						8/8-	21/-21	1/-1	3/-3	820/-820	-12/12	6/6-	44/-44	-13/13	3/-3	4/-4	9/9-	-4/4	8-/8	5/-5
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Right	79540 79529	79525	1047	1489	170	525	104614	1048	1491	1492	34855	34858	34861	76619	79601	76803	78464	69134	96769	69172	20767	88805	92632	101763	102686	107352	107551	19578	30525	30595	35899	52992	56214	61134	6201
Left	79538	79523	1046	1488	162	524	104613	1047	1490	1491	34852	34856	34858	76617	00962	76801	78462	69132	69294	69170	99/05	88783	92630	101759	101865	107351	107550	19533	30524	30591	35894	52991	56213	61125	6195
Marker Name	473729 473730	473731	473912	473913	473914	473915	474246	474247	474248	474249	474250	474251	474252	474253	474254	466999	467791	469419	469420	469421	470303	470304	471343	470130	470131	470132	470133	470134	470135	470136	470137	470138	470139	470140	470141
BAC Name	F5D14 F5D14	F5D14	F5D14	F5D14	F5D14	F5D14	F5D14	F5D14	F5A13	F2J6																									
BAC Chromosome Length	127462 127462	127462	127462	127462	127462	127462	127462	127462	127462	127462	127462	127462	127462	127462	127462	94618	94618	94618	94618	94618	94618	94618	94618	108061	190801	108061	108061	108061	108061	108061	108061	108061	108061	108061	108061
	AC007767 1 AC007767 1	AC007767 1	AC007767 1	AC007767 1	AC007767 1	AC007767 1	AC007767 1	AC007767 1	AC008046 1	AC009526 1																									
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	Right	68456	71329	72096	81241	72121	73969	4109	80466	83097	91920	98455	56240	56361	56248	45987	100539	1256	118609	121799	122815	16416	1762	19724	20647	27138	3579	39510	82738	85179	9881	55975	52760	52864	52972	52779	52744
	Left	68412	71322	72080	72103	72105	73956	7708	80459	83084	91873	98451	56238	56359	56246	45985	100537	1185	118608	121798	122811	16412	1761	19720	19826	27106	2762	39500	82675	85178	9834	55972	52758	52862	52970	52777	52742
Marker	Name	470142	470143	470144	470145	470146	470147	470148	470149	470150	470151	470152	467834	467835	467836	468042	468838	470073	470074	470075	470076	470077	470078	470079	470080	470081	470082	470083	470084	470085	470086	471276	471640	471641	471642	471643	471644
BAC	Name	F2J6	F2J6	F2J6	F216	F2J6	F28H19	F28H19	F28H19	F28H19	F28H19	F28H19	F28H19	F28H19	F28H19	F28H19	F28H19	F28H19	F28H19	F28H19	F28H19	F28H19	F28H19	F28H19	F28H19	F28H19	F27F5	F27F5	F27F5	F27F5	F27F5						
BAC	Chromosome Length	108061	108061	108061	108061	108061	108061	108061	108061	108061	108061	108061	131692	131692	131692	131692	131692	131692	131692	131692	131692	131692	131692	131692	131692	131692	131692	131692	131692	131692	131692	131692	137336	137336	137336	137336	137336
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	Sea id	AC009526	AC009526	AC009526	AC009526	AC009526	AC009526	AC009526	AC009526	AC009526	AC009526	AC009526	AC006423	AC006423	AC006423	AC006423	AC006423	AC006423	AC006423	AC006423	AC006423	AC006423	AC006423	AC006423	AC006423	AC006423	AC006423	AC006423	AC006423	AC006423	AC006423	AC006423	AC007915	AC007915	AC007915	AC007915	AC007915
Seg	unu kac	20	20	20	20	20	20	20	20	20	20	20	21	21	21	21	21	21	21	21	21	21	21	21	21	21	21	21	21	21	21	21	22	22	22	22	22

SNP Base Columbia/	Landsberg	C/I G/A	T/C	A/G	A/G	T/C	G/A	A/C	G/A	T/C	T/C	A/G	A/G	C/T	G/T	CT	G/A	T/A	G/A	C/A	T/C	D/L	C/T	C/A	C/A	A/C	A/C	A/G	A/G	G/T	G/C	A/G	G/A	G/A	G/A	T/C
Indel Size Columbia/	Landsberg																																			
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	Type	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP
	Right	51110	50980	51115	123658	122872	111586	111536	84991	85993	84910	85889	84834	85932	85991	85215	81733	79730	81384	81090	79775	81186	80939	77190	76656	76633	77508	77310	26800	76625	107289	105560	102965	102992	103060	102961
	Left	51108	50978	51113	123656	122870	111584	111534	84989	85991	84908	85887	84832	85930	82989	85213	81731	79728	81382	81088	79773	81184	80937	77188	76654	76631	77506	77308	86191	76623	107287	105558	102963	102990	103058	102959
Marker	Name	471722	471723	471724	471808	471833	472413	472414	472494	472495	472496	472497	472498	472499	472500	472501	472509	472510	472511	472512	472513	472514	472515	472516	472517	472518	472519	472520	472521	472522	472573	472717	472718	472719	472720	472721
BAC	Name	F27F5	F27F5	F27F5	F27F5	F27F5	F27F5	F27F5	F27F5	F27F5	F27F5	F27F5	F27F5	F27F5	F27F5	F27F5	F27F5	F27F5	F27F5	F27F5	F27F5	F27F5	F27F5	F27F5	F27F5	F27F5	F27F5	F27F5	F27F5	F27F5	F27F5	F27F5	F27F5	F27F5	F27F5	F27F5
BAC	Chromosome Length	1 137336	1 137336	1 137336	1 137336	1 137336	1 137336	1 137336	1 137336	1 137336	1 137336	1 137336	1 137336	1 137336	1 137336	1 137336	1 137336	1 137336	1 137336	1 137336	1 137336	1 137336	1 137336	1 137336	1 137336	1 137336	1 137336	1 137336	1 137336	1 137336	1 137336	1 137336	1 137336	1 137336	1 137336	1 137336
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SNP Base Columbia/ Landsberg G/C C/T C/T T/G T/G T/G C/A	17.G G/A G/C G/C C/A G/C
Indel Size Columbia/ Landsberg -1/1 -2/2 1/-1 1/-1 1/-7 -1/1	16/-16 3/-3 -3/3 10477/- 10477 -4/4 10/-10 3/-3 19/-19
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Type SNP SNP SNP SNP SNP SNP SNP SNP SNP SNP SNP SNP SNP	S S S S S S S S S S S S S S S S S S S
Right 103115 102870 103024 113951 92993 93113 102389 101925 101938 122428 122428 122428 122428 34758	34634 34939 71807 49217 49070 54638 55430 18021 43189 43025 42909 30823 30998 31038 47227 39304 39658 49083 51307
Left 103113 102868 103022 113949 92991 93111 101924 101937 102271 122426 124140 89612	34632 34632 34997 71805 49215 49068 55428 18019 43187 43023 42907 30806 30994 31037 36749 39303 39647 49079 5731
Marker Name 472722 472723 472724 473639 473659 473659 474161 474162 474163 474164 474165 474166	467029 467029 467030 467898 467899 468214 468411 468486 470956 470956 470959 470960 470963 470963 470963
BAC Name F27F5	T3F24 T3F24 T3F24 T3F24 T3F24 T3F24 T3F24 T3F24 T3F24 T3F24 T3F24 T3F24 T3F24 T3F24
BAC ome Length 137336 137336 137336 137336 137336 137336 137336 137336 137336 137336 137336	77424 77424 77424 77424 77424 77424 77424 77424 77424 77424 77424 77424
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SNP Base Columbia/ Landsberg	A/G C/A C/G C/G A/C	T/C A/G G/A T/C C/T T/A T/C
Indel Size Columbia/ Landsberg 292/-292 37/-37 -8/8 3/-3 3/-3 2/-2	i i	2577/-2577 4/-4 10/-10 -9/9 -4/4 25084- 25084- 25084 25084- 25084 2508- 15/15 -15/15 -17/1
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Type IND IND IND IND IND	SNP SNP SNP SNP SNP SNP SNP SNP SNP SNP	
Right 65066 66894 77010 7754 7758 34599	50797 50687 15355 75126 75663 84324 84329	84527 84426 1756 1684 1657 9331 202 73654 73697 19905 18641 25126 37534 68815 48161 71870 72308 15403 47510
Left 64773 66856 77009 7750 7754 34596	50795 50685 15353 75124 75661 84322 84327	84525 84424 1754 1682 1682 1655 9329 200 73652 73695 17327 18636 25115 35204 37533 43730 15400 47564 72307
Marker Name 470965 470966 470967 470968 471579	471380 467168 467169 467489 467935 468257 468258	468259 468359 468330 468331 468344 469239 469461 470909 470910 470911 470914 470915 470915 470915 470916 471561
BAC Name T3F24 T3F24 T3F24 T3F24 T3F24 T3F24	13F24 12E6 12E6 12E6 12E6 12E6 12E6	1256 1256 1256 1256 1256 1256 1256 1256
BAC Chromosome Length 1 77424 1 77424 1 77424 1 77424 1 77424	97154 97154 97154 97154 97154 97154 97154	97154 97154 97154 97154 97154 97154 97154 97154 97154 97154 97154 97154 97154 97154
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SNP Base Columbia/ Landsberg C/A A/G C/T A/T A/T A/T A/T A/T C/A G/C C/G G/C C/A T/C C/A C/A C/A C/A C/A C/A C/A C/A C/A C
Indel Size Columbia/ Landsberg -1/1
Method
SNP
Right 94140 111880 76928 76737 87314 87314 87228 87439 18915 113710 113701 113896 13163 12917 47777 68600 67150 66944 66947 66947 66947 66982 66114 39380 36926 100067 100065 100655 100655
Left 94139 111878 73939 76926 76735 87312 87312 87226 87437 18913 113731 113731 113739 113894 113894 113899 113894 113899 113894 13161 12762 12915 47775 66945 66945 66945 66945 66945 100659 100653 100653 100663 100633
Marker Name 471564 466884 466884 467011 467012 467053 467053 467053 467250 467251 467252 467253 467253 467253 467351 467351 468171 468173 468177 468177 468177 468178 468177 468177 468177 468178 468177 468177 468886 468889 468889
BAC Name T2E6 F27J15
BAC Chromosome Length 1 19091 1 19091
Seq id AC012463 AC012463 AC012463 AC016041
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Indel Size	Landsberg														4/-4	9-/9	4/-4	13/-13	9-/9	ê/8-	-36/36	4/-4	4/-4	-3/3	-4/4	6/6-	12/-12	-21/21	-21/21	-5/5	3/-3	206/-206	30/-30	-15/15	-4/4	-12/12
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	Right	100552	100129	101342	85006	85047	7165	7203	7273	118145	118569	115575	117818	115362	100576	10274	1209	14576	38058	41841	41990	42412	42417	47911	51562	55711	56647	57120	57125	58287	5837	59093	59514	60541	79671	81311
	Left 100550	100539	100127	101340	85004	85045	7163	7201	7271	118143	118567	115573	117816	115360	100571	10267	1204	14562	38051	41840	41989	42407	42412	47910	51561	55710	56634	57119	57124	58286	5833	58886	59483	60540	02962	81310
Marker	Name 468804	468895	468896	468897	469014	469015	469276	469277	469278	469368	469369	469370	469371	469372	470047	470048	470049	470050	470051	470052	470053	470054	470055	470056	470057	470058	470059	470060	470061	470062	470063	470064	470065	470066	470067	470068
BAC	Name F27115	F27J15	F27J15	F27J15	F27J15	F27J15	F27J15	F27J15	F27J15	F27J15	F27J15	F27J15	F27J15	F27J15	F27J15	F27J15	F27J15	F27J15	F27J15	F27J15	F27J15	F27J15	F27J15	F27J15	F27J15	F27J15	F27J15	F27J15	F27J15	F27J15	F27J15	F27J15	F27J15	F27J15	F27J15	F27J15
BAC	Chromosome Length	119091	119091	119091	119091	119091	119091	119091	119091	119091	119091	119091	119091	119091	119091	119091	119091	119091	119091	119091	119091	119091	119091	119091	119091	119091	119091	119091	119091	119091	119091	119091	119091	119091	119091	119091
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Right 83562 89225 96468 96811 100509 100576 100580 100586 100685 12774 28853	32198 67648 67528 90894 23798 10925 10893	10893 10893 10997 10949 33688 92291 88505 88503 88503 87982 34623 7999 8119 37022 37185 37211
Left 83553 89221 96467 96785 100574 100578 100578 100583 12773	32196 67646 67526 90892 23796 10923	10891 10995 10995 10947 33686 92289 88591 87980 88605 28123 34621 7997 8117 37685 37020 37183
Marker Name 470069 470070 470071 471270 471271 471273 471273 471273	466801 466815 466816 467034 467303 467343	467345 467345 467347 467643 467643 467670 467670 467671 468126 468567 468567 468571 468571 468571
BAC Name F27J15	F14322 F14322 F14322 F14322 F14322 F14322	F14522 F14522 F14522 F14522 F14522 F14522 F14522 F14522 F14522 F14522 F14522 F14522 F14522 F14522 F14522
BAC me Length 119091 119091 119091 119091 119091 119091 119091 119091	104679 104679 104679 104679 104679 104679	104679 104679 104679 104679 104679 104679 104679 104679 104679 104679 104679 104679
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Right 63979 70638 78377 84087 84168 85005 89459	120689 120987 54745 122380 124790 124899	124910 125052 124746 124805 129005 67031 75160 74648 74208 74602 74602 74601 74585 109435 109264 109264	113795 117256 133525 133547
Left 63978 70634 78373 84030 84159 85004 89454	120687 120985 54743 122378 124788 124897	124908 125050 124803 124803 129003 67029 75158 74646 74206 74000 74059 74583 109433 109260 109260	113790 117227 133520 133546
Marker Name 470123 470124 470125 470126 470127 470128	466883 466883 467143 467691 467802 467803	467805 467806 467807 467808 469408 469412 469414 469415 469416 469444 469444 469446 469447 469448	469985 469986 469987 469988
BAC Name F2J10 F2J10 F2J10 F2J10 F2J10 F2J10	F25P12 F25P12 F25P12 F25P12 F25P12 F25P12	F25P12 F25P12 F25P12 F25P12 F25P12 F25P12 F25P12 F25P12 F25P12 F25P12 F25P12 F25P12 F25P12	F25P12 F25P12 F25P12 F25P12
BAC Chromosome Length 1 91720 1 91720 1 91720 1 91720 1 91720	141753 141753 141753 141753 141753 141753	141753 141753 141753 141753 141753 141753 141753 141753 141753 141753 141753 141753 141753	141753 141753 141753 141753
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Type Can and C	SNP SNP SNP SNP SNP	SNP SNP SNP SNP SNP SNP SNP SNP SNP SNP
Right 138261 140783 57206 6700 67556 7731 96672 96864 96944 109272 124820 66417 74588	84205 84146 84041 84245 84227	84091 83985 20830 20982 3827 3586 37234 43441 43547 4358 43682 43682 39565 39565 39565 39505
Left 138260 140782 57191 6686 67547 7730 96664 96845 96935 109270 124818 66416 74587	84203 84144 84039 84243 84225	84089 83983 20828 20820 3825 3584 37232 43439 43545 43456 43680 43744 39563 39563 39503 34987
Marker Name 469989 469990 469991 469993 469995 469996 469996 471242 471244 471244	471820 471821 471822 471823	471825 471826 472044 472045 472045 472311 472486 472486 472486 472487 472487 472487 472582 472583 472583 472583
BAC Name F25P12	F19C14 F19C14 F19C14 F19C14 F19C14	F19C14 F19C14 F19C14 F19C14 F19C14 F19C14 F19C14 F19C14 F19C14 F19C14 F19C14 F19C14 F19C14
BAC Chromosome Length 1 141753 1 141753 1 141753 1 141753 1 141753 1 141753 1 141753 1 141753 1 141753 1 141753 1 141753 1 141753 1 141753	86014 86014 86014 86014 86014	86014 86014 86014 86014 86014 86014 86014 86014 86014 86014 86014 86014 86014 86014 86014
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Right 27134 27078 27078 27036 27137 11879 12054 13399 13400 14168 13413 8857 9349 8907 41236 25562 25427 26920 3556 43526 58031 29772 29811 11284 111524 11173 11173	11342 12927 10728 11611 61517 61332 61161
Left 27132 27036 27034 27135 11877 12052 13398 14166 13411 8855 9347 8905 41234 25560 25425 26917 3554 43524 58030 29770 29809 38051 11522 11639 11152	112925 12925 10726 11609 61515 61330 61159
Marker Name 473104 473105 473106 473106 473125 473291 473292 473293 473293 473293 473293 473294 473294 473506 473506 473506 473508 47360 473508 47360 47360 47360 47360 47360 47360 47360 472067 472363 472365 472365	472369 472370 472371 472529 472530
BAC Name F19C14 F19C17 F16P17 F16P17 F16P17 F16P17 F16P17 F16P17	F16P17 F16P17 F16P17 F16P17 F16P17
BAC Chromosome Length 86014 86014 86014 86014 1 86014	98412 98412 98412 98412 98412 98412
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SNP Base	Landsberg	A/G	A/G	C/G	G/A	T/A	C/A	1/A	7/G	D/1 D/T	C/T	G/A	T/A	A/G	) L	T/A	A/T	. T.	G/A	T/C	D/L	A/G	7/U	G/T	A/T	I !										
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	Right	61729	61/02	40028	49020	49304	7164	49022	49322	49754	37471	36161	74649	74442	74439	17999	17934	17976	51642	50684	50377	51703	50356	51562	14200	3110	3247	3441	5962	12830	12833	12837	12896	18049	49296	49421
	Left	61727	01/00	01230	49020	49502	49665	49020	49320	49752	37469	36159	74647	74440	74437	17997	17932	17974	51640	50682	50375	51701	50354	51560	14198	2593	3246	3437	5953	12828	12830	12833	12895	18047	49295	49418
Marker	Name	4/2532	472533	472916	472017	472918	472919	472920	472921	472922	472945	473116	473158	473159	473160	473204	473205	473206	473755	473756	473757	473758	473759	473760	473797	473842	473843	473844	473845	474075	474076	474077	474078	474079	474080	474081
BAC	Name	F10F1/	F10F1/ F16P17	F101.1/ F16P17	F16P17	F16P17	F16P17	F16P17	F16P17	F16P17	F16P17	F16P17	F16P17	F16P17	F16P17	F16P17	F16P17	F16P17	F16P17	F16P17	F16P17	F16P17	F16P17	F16P17	F16P17	F16P17	F16P17	F16P17	F16P17	F16P17	F16P17	F16P17	F16P17	F16P17	F16P17	F16P17
BAC	Chromosome Length	98412	98412	98412	98412	98412	98412	98412	98412	98412	98412	98412	98412	98412	98412	98412	98412	98412	98412	98412	98412	98412	98412	98412	98412	98412	98412	98412	98412	98412	98412	98412	98412	98412	98412	98412
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SNP Base Columbia/ Landsberg	G/A 1/A 1/C 1/C 1/C 1/C		G/A C/T G/A T/A T/A T/C G/C
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Right 61245 61646 88619 88813	31523 50730 50888 49002 48870 57812 56711 57764	32990 34332 38275 65445 65509 66563 66706 7904 89549 9167 92714 92716 49978 86850	47763 48638 37406 37223 57268 56919 57033
Left 61244 61644 88618 88812	31521 50728 50886 49000 48868 57810 56709 57762 56334	32989 33989 33286 65444 65508 66702 7887 89548 9166 92713 92713 86849	47761 48636 37404 37221 57266 56917 57031
Marker Name 474082 474083 474084	467124 467204 467205 467207 467601 467603 467603	469695 469696 469697 469699 469700 469701 469703 469704 469705 469705 471155	467719 467720 467867 467868 468651 468652 468653
BAC Name F16P17 F16P17 F16P17	F16M19 F16M19 F16M19 F16M19 F16M19 F16M19 F16M19 F16M19	F16M19 F16M19 F16M19 F16M19 F16M19 F16M19 F16M19 F16M19 F16M19 F16M19	F2K11 F2K11 F2K11 F2K11 F2K11 F2K11 F2K11
BAC Chromosome Length 1 98412 1 98412 1 98412	100512 100512 100512 100512 100512 100512 100512 100512	100512 100512 100512 100512 100512 100512 100512 100512 100512 100512 100512 100512 100512	100867 100867 100867 100867 100867 100867
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Indel Size Columbia/	Landsberg																											3/-3	-64/64	-5/5	-3/3	-11/11	-3/3	-3/3	11/-11	58/-58	3/-3
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	Right	5/263	56921	5/182	53628	53845	23767	53446	94588	95568	94669	96046	20457	20421	64811	64851	63884	50302	18894	15893	25494	73361	70722	70781	69580	68969	69425	12925	26520	34636	349	363	36334	38272	40160	54650	69009
	Left	27.201	56919	57.180	33626	53843	53765	53444	94586	95566	94667	96044	20455	20419	64809	64849	63882	50300	18892	15891	25492	73359	70720	67101	69578	28969	69423	12921	26519	34635	348	362	36333	38271	40148	54591	9009
Marker	Name	408033	408050	468637	408/91	468/92	468793	468794	468855	468856	468857	468858	468960	468961	468986	468987	468988	469078	469128	469129	469130	469240	469241	469242	469243	469244	469245	470153	470154	470155	470156	470157	470158	470159	470160	470161	470162
BAC	Name	F277.1	F2K11	FZKII	FZRII	F2K11	F2K11	F2K11	F2K11	F2K11	F2K11	F2K11	F2K11	F2K11	F2K11	F2K11	F2K11	F2K11	F2K11	F2K11	F2K11	F2K11	F2K11	F2K11	F2K11	F2K11	F2K11	F2K11	F2K11	F2K11	F2K11	F2K11	F2K11	F2K11	F2K11	F2K11	F2K11
BAC	Chromosome Length	100807	100867	100867	100867	100007	100867	100867	100867	100867	100867	100867	100867	100867	100867	100867	100867	100867	100867	100867	100867	100867	100867	100867	100867	100867	100867	100867	100867	100867	100867	100867	100867	100867	100867	100867	100867
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SNP Base	Londoborg	Lanusocie						G/A	C/T	T/A	T/C	C/A	G/A	G/A	G/A	T/C	G/C	G/C	A/G	A/G	A/G	C/G	A/G	C/T	G/T	AVT	C/T										
Indel Size	I andshara	-60/60	63/-63	1/-1	-1/1	1/-1	-1/1																					-4/4	-4/4	9-/9	9-/9	8-/8	5/-5	-5/5	6-/6	8-/8	8-/8
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	Rioht	69151	8776	100612	48518	69622	66869	84930	86137	74402	74436	72319	71217	71218	71536	72448	71577	71602	73078	71390	71539	71604	71667	71491	71562	71664	71676	2833	30697	37385	41834	45499	49480	49540	50990	69489	69535
	Left	69150	8712	100610	48517	69620	86869	84928	86135	74400	74434	72317	71215	71216	71534	72446	71575	71600	73076	71388	71537	71602	71665	71489	71560	71662	71674	2832	30696	37378	41827	45490	49474	49539	50980	69480	69526
Marker	Name	470163	470164	471297	471298	471299	471300	467528	468373	469003	469004	469373	469374	469375	469376	469377	469378	469379	469380	469381	469382	469383	469384	469385	469386	469387	469388	469562	469563	469564	469565	469566	469567	469568	469569	469570	469571
BAC	Name	F2K11	F2K11	F2K11	F2K11	F2K11	F2K11	F12B7	F12B7	F12B7	F12B7	F12B7	F12B7	F12B7	F12B7	F12B7	F12B7	F12B7	F12B7	F12B7	F12B7	F12B7	F12B7	F12B7	F12B7	F12B7	F12B7	F12B7	F12B7	F12B7							
BAC	Chromosome Length	100867	100867	100867	100867	100867	100867	91530	91530	91530	91530	91530	91530	91530	91530	91530	91530	91530	91530	91530	91530	91530	91530	91530	91530	91530	91530	91530	91530	91530	91530	91530	91530	91530	91530	91530	91530
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	Seq id	AC008047	AC008047	AC008047	AC008047	AC008047	AC008047	AC011020	AC011020	AC011020	AC011020	AC011020	AC011020	AC011020	AC011020	AC011020	AC011020	AC011020	AC011020	AC011020	AC011020	AC011020	AC011020	AC011020	AC011020	AC011020	AC011020	AC011020	AC011020	AC011020							
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SNP Base Columbia/ Landsberg	C/T C/A T/G A/T	AT G/A C/T C/T C/T A/T C/T C/T C/T	A/T G/T T/A T/A
Indel Size Columbia/ Landsberg 3/-3 3/-3 -9/9 -6/6 7/-7 5/-5 3/-3 8/-8 -6/6 -62/62	-1/1		
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Right 70017 70032 70041 70140 70927 71130 71147 75873 81921 85194 72310 744330	74484 86418 97659 63743 7655 8486	28253 28253 28207 84005 76307 94578 94759 94874 24127 24933 48611 47822 47959	96394 60023 35445 35393
Left 70013 70028 70040 70139 71124 71143 71143 71143 71143 7143 7143 81920 85193 74328	74463 86416 97657 63741 7653 8484	28251 28205 84003 76305 94576 94757 94872 24931 48609 47957 96135	96392 60021 35443 35391
Marker Name 469572 469573 469574 469575 469577 469578 469580 469581 471096 471096	46879 466879 467295 467312 467313	467730 467731 467731 468014 468022 468023 468040 468040 468361 468362	468435 468445 468454 468455
BAC Name F12B7 F12B7 F12B7 F12B7 F12B7 F12B7 F12B7 F12B7 F12B7 F12B7 F12B7	F14023 F14023 F14023 F14023 F14023	F14023 F14023 F14023 F14023 F14023 F14023 F14023 F14023 F14023 F14023	F14023 F14023 F14023 F14023
BAC Chromosome Length 1 91530 1 91530 1 91530 1 91530 1 91530 1 91530 1 91530 1 91530 1 91530 1 91530	98471 98471 98471 98471 98471	98471 98471 98471 98471 98471 98471 98471 98471 98471	98471 98471 98471 98471
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SNP Base		C/A	7/C	A/G	A/T	AT	CT	T/A	A/C	T/G																						T/C	T/A	T/A	G/A	G/A
Indel Size Columbia/	Landsberg										-5/5	10/-10	9/9-	9/9-	4/-4	11-7	3/-3	3/-3	L/L-	-40/40	1/-1	-1/1	1/-1	1/-1	1/-1	1/-1	9-/9	-1/1	1/-1	-1/1	1/-1					
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	Type	JANS:	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	ONI	IND	QNI	IND	QNI	ONI	IND	ONI	IND	ONI	IND	ONI	ONI	ONI	IND	ONI	ONI	IND	ONI	ONI	IND	SNP	SNP	SNP	SNP	SNP
	Right	35252	35972	35884	35916	9821	22869	62037	62483	62212	12485	22321	22364	22365	340	37045	52883	77537	81303	88903	24679	34941	35945	35973	36240	37045	37057	37167	83764	86385	06096	20972	64108	65650	65610	65472
	1500 Jeft	35250	35970	35882	35914	9819	69875	62035	62481	62210	12484	22310	22363	22364	335	37037	52879	77533	81302	88902	24677	34940	35943	35971	36238	37043	37050	37166	83762	86384	88096	20970	64106	65648	80959	65470
Marker	Name 468456	468457	468458	468459	468460	468962	469089	469119	469120	469121	469648	469649	469650	469651	469652	469653	469654	469655	469656	469657	471136	471137	471138	471139	471140	471141	471142	471143	471144	471145	471146	466850	467077	467078	467079	467080
BAC	Name F14023	F14023	F14023	F14023	F14023	F14023	F14023	F14023	F14023	F14023	F14023	F14023	F14023	F14023	F14023	F14023	F14023	F14023	F14023	F14023	F14023	F14023	F14023	F14023	F14023	F14023	F14023	F14023	F14023	F14023	F14023	F25P22	F25P22	F25P22	F25P22	F25P22
BAC	Chromosome Length	98471	98471	98471	98471	98471	98471	98471	98471	98471	98471	98471	98471	98471	98471	98471	98471	98471	98471	98471	98471	98471	98471	98471	98471	98471	98471	98471	98471	98471	98471	105937	105937	105937	105937	105937
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SNP Base	Landsberg	) [ [	A/G	AT.	CT	G/A	T/C	T/A	AT	A/T	G/A	T/A	T/C	C/G	T/C	CT	G/T	C/C	T/C	T/C	A/G	A/G	A/G	T/G	I/G	C/T	A/T	G/T	A/T	C/T	C/T	C/T	T/C	T/C	G/A	C/A
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	Right	65656	27556	102782	71236	19882	18549	78543	78598	78547	31153	31116	31079	30956	13934	13936	58105	63575	63584	62878	63563	63587	63590	63593	63609	63085	63538	63553	63562	62824	62830	62854	52560	53370	99553	99634
	Left	65654	27554	102780	71234	19880	18547	78541	78596	78545	31151	31114	31077	30954	13932	13934	58103	63573	63582	62876	63561	63585	63588	63591	63607	63083	63536	63551	63560	62822	62828	62852	52558	53368	99551	99632
Marker	Name	467082	467338	467418	467721	467776	467777	467883	467884	467885	468031	468032	468033	468034	468369	468370	468380	468622	468623	468624	468625	468626	468627	468628	468629	468630	468631	468632	468633	468634	468635	468636	468639	468640	469254	469255
BAC	Name	F25P22	F25P22	F25P22	F25P22	F25P22	F25P22	F25P22	F25P22	F25P22	F25P22	F25P22	F25P22	F25P22	F25P22	F25P22	F25P22	F25P22	F25P22	F25P22	F25P22	F25P22	F25P22	F25P22	F25P22	F25P22	F25P22	F25P22	F25P22	F25P22	F25P22	F25P22	F25P22	F25P22	F25P22	F25P22
BAC	Chromosome Length	105937	105937	105937	105937	105937	105937	105937	105937	105937	105937	105937	105937	105937	105937	105937	105937	105937	105937	105937	105937	105937	105937	105937	105937	105937	105937	105937	105937	105937	105937	105937	105937	105937	105937	105937
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	Seq id AC012679	AC012679	AC012679	AC012679	AC012679	AC012679	AC012679	AC012679	AC012679	AC012679	AC012679	AC012679	AC012679	AC012679	AC012679	AC012679	AC012679	AC012679	AC012679	AC012679	AC012679	AC012679	AC012679	AC012679	AC012679	AC012679	AC012679	AC012679	AC012679	AC012679	AC012679	AC012679	AC012679	AC012679	AC012679	AC012679
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SNP Base Columbia/	Landsberg T/G	)																						T/A	AC	G/C	T/G	C/A	T/A	T/A	T/A	T/A	T/C	CT	G/T	A/T
Indel Size Columbia/	Landsberg	6/6-	5/-5	-4/4	3/-3	13/-13	L-1/L	17/-17	4/-4	9/9-	14/-14	10/-10	3/-3	1/-1	1/-1	1/-1	-2/2	1/-1	1/-1	-1/1	-1/1	-2/2	-1/1													
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	Right 74002	11135	26268	26470	28236	2979	31322	35124	35194	36344	3830	3986	86730	52252	56217	56224	57506	61334	63607	65585	65586	65646	98602	75428	88209	86763	88850	6096	15181	92581	93849	90812	06606	91009	92909	38949
	Left 74000	11134	26262	26469	28232	2965	31314	35106	35189	36343	3815	3975	86726	52250	56215	56222	57505	61332	63605	65584	65585	65645	70985	75426	88207	86761	88848	2096	15179	92579	93847	90810	88606	20016	60674	38947
Marker	Name 469354	469998	469999	470000	470001	470002	470003	470004	470005	470006	470007	470008	470009	471247	471248	471249	471250	471251	471252	471253	471254	471255	471256	472068	472221	472222	472223	472276	472932	473043	473044	473118	473119	473120	473153	473485
BAC	Name F25P22	F25P22	F25P22	F25P22	F25P22	F25P22	F25P22	F25P22	F25P22	F25P22	F25P22	F25P22	FIB16	F1B16																						
BAC	Chromosome Length 1 105937	105937	105937	105937	105937	105937	105937	105937	105937	105937	105937	105937	105937	105937	105937	105937	105937	105937	105937	105937	105937	105937	105937	100685	100685	100685	100685	100685	100685	100685	100685	100685	100685	100685	100685	100685
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	Seq 1d AC012679	AC012679	AC012679	AC012679	AC012679	AC012679	AC012679	AC012679	AC012679	AC012679	AC012679	AC012679	AC023754																							
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SNP Base Columbia/ Landsberg G/A C/A T/G	7,C C/T C/G C/G C/G C/G C/G C/G C/G C/G C/G C/G
Indel Size Columbia/ Landsberg -6/6 25/-25 4/-4 -6/6 -1/1 1/-1 -1/1 -1/1 -6/6	7-17
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Right 36320 10726 10446 10792 21509 381 5590 8379 10450 13171 23714 77984 8258 8378	25603 25603 25603 25305 47642 6048 5885 6515 6299 46196 33418 37041 32931
Left 36318 10724 10444 10790 21508 355 5585 8378 10449 13169 23713 77983 8257 8377	25601 25601 25601 25303 47640 6046 5883 6297 46194 33680 33416 37039 32929 32929 32929 32929 32929 32929 32929 32929 32929 32929 32929 32929
Marker Name 473492 473808 473809 473847 473847 473849 473849 474096 474097 474099 474100	4/4102 466926 466928 467348 467348 467570 467571 468501 468502 468503 468504 468504 468504 468504 468504 468506 468509 468509 468509
BAC Name FIB16	7.15.10 7.23.51.8 7.23.51.8 7.23.51.8 7.23.51.8 7.23.51.8 7.23.51.8 7.23.51.8 7.23.51.8 7.23.51.8 7.23.51.8 7.23.51.8 7.23.51.8 7.23.51.8 7.23.51.8 7.23.51.8 7.23.51.8
BAC Chromosome Length 1 100685 1 100685	97554 97554 97554 97554 97554 97554 97554 97554 97554 97554 97554 97554 97554 97554
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SNP Base Columbia/ Landsberg T/C A/G T/A T/A T/C A/C T/A	T/C
Landsberg Columbia/ Landsberg 23/-23 11/-11 -13/13 35/-35 4/-4 9/-9 -5/5 -11/11 34/-34 13/-13 4/-4 7/-7 10/-10 4/-4 -2/2 1/-1 -2/2 1/-1 -2/2	
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Right 45241 45241 45243 30189 43534 10809 11505 12392 11941 10805 12331 13650 14395 1078 17122 1874 1890 27073 37250 45957 46150 60904 63580 664493 79581 13696 44585 44773 46151 38437	11/40
Left 45239 45221 30187 43532 10807 11503 12569 12390 11939 10803 12329 13648 14393 1077 17098 1862 1889 27037 37245 46149 60864 60864 60869 63572 63672 63672 63463 13695 44584 44771	11/44
Marker Name 468602 468603 468902 468902 469291 469291 469292 469295 469295 469295 469295 469297 469296 469297 469297 469297 469297 469297 469297 470835 470839 470849 470849 470849 470848 470849 470848 470849 470849 470849	40/0/3
BAC Name T23E18	OTTATOT
BAC Chromosome Length 97554 1 97554 1 97554 1 97554 1 97554 1 97554 1 97554 1 97554 1 97554 1 97554 1 97554 97554 97554 97554 97554 97554 97554 97554 97554 97554 97554 97554 97554 97554	000001
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SNP Base	Landsberg	G/A	T/C	T/C	A/T	C/T	C/T	A/T	T/A	T/C	T/A	G/C	G/C	T/C	G/A	A/G	A/G	G/A	T/C	T/G	A/C	C/G	A/T	G/T	C/T	G/C	T/C	G/T	A/T	T/C	A/C	A/T	A/G	G/C	T/C	J/G	G/T
Indel Size Columbia/	Landsberg																																				
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	Type	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP
	Right	66285	68562	66803	009/9	66362	66364	69063	34206	33197	30364	31362	30032	30033	55198	56526	102182	3986	4226	91042	8985	27115	96264	95578	27956	83257	83578	83428	83060	82861	82249	82122	43076	17796	21925	21686	18085
	Left	66283	09589	66801	67598	09899	66362	69091	34204	33195	30362	31360	30030	30031	55196	56524	102180	3984	4224	91040	8983	27113	96262	92576	27954	83255	83576	83426	83058	82859	82247	82120	43074	17794	21923	21684	18083
Marker	Name	467587	467588	467589	467590	467591	467592	467593	467621	467622	467623	467624	467625	467626	467629	467630	467680	467726	467727	467736	467781	467906	467939	467940	467978	467997	467998	467999	468000	468001	468002	468003	468093	468094	468095	468096	468097
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BAC	Chromosome Length	103353	103353	103353	103353	103353	103353	103353	103353	103353	103353	103353	103353	103353	103353	103353	103353	103353	103353	103353	103353	103353	103353	103353	103353	103353	103353	103353	103353	103353	103353	103353	103353	103353	103353	103353	103353
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Sed	unu	38	38	38	38	38	38	38	38	38	38	38	38	38	38	38	38	38	38	38	38	38	38	38	38	38	38	38	38	38	38	38	38	38	38	38	38

e SNP Base	_	A/T	T/C	G/A	G/C	T/C	T/C	A/G	A/T	A/T	A/T	A/C	T/A	T/C	T/C	A/G	T/G	C/G	C/A	G/A	T/C	C/T	A/T	C/T	A/T	T/A	G/A	G/C	G/C	A/G	A/G	G/T	A/C	T/A			
Indel Size Columbia/	Landsberg																																		8-/8	9-/9	-3/3
	Method	-	-		_	1	1	1	1	1	-		1	T	1	1	Т	1	1	1	1	7		<del></del>	-		1	1	1	1	1	I	1		2	2	7
	Type	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	ONI	IND	ONI																						
	Right	18413	87460	86668	89107	90280	90030	90037	88833	88507	88506	63085	79832	79831	77340	77583	76576	78884	85876	85663	84446	84592	35171	35231	49340	50703	51172	50591	50946	50609	51102	51103	60582	41572	102037	11827	13932
	Left	18411	87458	89991	89105	90278	90028	90035	88831	88505	88504	63083	79830	79829	77338	77581	76574	78882	85874	85661	84444	84590	35169	35229	49338	50701	51170	50589	50944	. 20905	51100	51101	60580	41570	102028	11820	13931
Marker	Name	468098	468184	468185	468186	468187	468188	468189	468190	468191	468192	468293	468474	468475	468476	468477	468478	468479	468658	468659	468660	468661	468724	468725	468770	468771	468772	468773	468774	468775	468776	468777	468901	469316	470980	470981	470982
BAC	Name	T5M16	T5M16	T5M16	T5M16	T5M16	T5M16	T5M16	T5M16	T5M16	T5M16	T5M16	T5M16	T5M16	T5M16																						
BAC	Chromosome Length	1 103353	1 103353	1 103353	1 103353	1 103353	1 103353	1 103353	1 103353	1 103353	1 103353	1 103353	1 103353	1 103353	1 103353	103353	103353	103353	103353	103353	103353	103353	103353	103353	103353	103353	103353	103353	103353	103353	103353	103353	103353	103353	103353	103353	103353
	Seq id	AC010704	AC010704	AC010704	AC010704	AC010704	AC010704	AC010704	AC010704	AC010704	AC010704	AC010704	AC010704	AC010704	AC010704																						
Seq	· E	38	38	38	38	38	38	38	38	38	38	38	38	38	38	38	38	38	38			38	38	38	38	38	38	•	•	•	•	38	38	•	Ì	,	38

SNP Base Columbia/ Landsberg	C/A A/G A/T G/A	C/T T/A G/A C/T A/G C/A C/A
Indel Size Columbia/ Landsberg -4/4 -7/7 15/-15 115/-15 -3/3 3/-3 3/-3 3/-3 3/-3 -3/3 -6/6 9/-9 4/-4 2/-2 1/-1 -1/1 -1/1 -1/1 -1/1 -1/1 -1/1		
Method 1 1 1 1 1 2 2 2 2 2 2 2 2 2 2 2 2 2 2 2		
	SNP SNP SNP SNP	SNP SNP SNP SNP SNP SNP SNP
Right 14122 25173 44320 45773 52948 53052 58441 58441 102041 102044 102046 2159 33940 3879 41666 78908 85690 88502 96502	72030 32908 32766 85675	74721 15190 41502 35483 19908 28268 28267 29263
Left 14121 25172 44304 45657 52947 53051 58447 78905 85670 99425 102040 102040 102044 2158 33939 3878 41664 78907 78908 88500 88500 88501	72028 32906 32764 85673	74719 15188 41500 35481 19906 28266 28265 29261
Marker Name 470983 470984 470985 470986 470986 470990 470991 471585 471585 471589 471599 471591 471591 471593 471594 471595 471595	467208 467716 467717 467986	468228 468446 468584 468769 468882 469471 469473
BAC Name TSM16	F3F9 F3F9 F3F9 F3F9	F3F9 F3F9 F3F9 F3F9 F3F9 F3F9
BAC Chromosome Length 1 103353	95771 95771 95771 95771	95771 95771 95771 95771 95771 95771
Chrom		
Seq id AC010704	AC013430 AC013430 AC013430 AC013430	AC013430 AC013430 AC013430 AC013430 AC013430 AC013430 AC013430
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SNP Base Columbia/ Landsberg A/G A/T C/T A/G A/G A/T A/G A/G	G/A 1/C A/T C/A G/T C/A
Indel Size Columbia/ Landsberg 22/-22 -4/4 7/-7 8/-8 7/-7 -4/4 -4/4 35/-35 63/-63 8/-8 -3/3 3/-3 6/-6 12/-12 -4/4 7/-7 -3/3 -3/3 -1/1 -1/1	10/-10
Method 1	2 2 11 11 11 11 11 1
	SNP SNP SNP SNP SNP SNP SNP SNP SNP SNP
Right 28811 28273 10189 15376 21175 21883 23789 23824 3255 38300 48455 5962 6705 74197 80170 80693 8183 17005 29338 32741 4802 4207 24876 9740	66113 66330 66365 64991 67513 67621 69930 15312 47343
Left 28809 28271 10166 15375 21167 21874 23823 3254 38264 48391 5953 6701 73998 74184 80169 80685 8181 8182 17004 29337 32738 4800 4205	9738 66111 66328 66363 64989 67511 67619 69928 15301
Marker Name 469474 469475 470236 470237 470238 470241 470242 470243 470244 470245 470245 470246 470246 470246 470253 471316 470253 471318 467130 467130	408443 468614 468615 468616 468617 468619 469288 469288
BAC Name F3F9 F3F9 F3F9 F3F9 F3F9 F3F9 F3F9 F3F	F20B17
BAC Chromosome Length  95771	90149 90149 90149 90149 90149 90149 90149
Seq id AC013430	AC010793 1 AC010793 1 AC010793 1 AC010793 1 AC010793 1 AC010793 1 AC010793 1 AC010793 1
Seq m m m m m m m m m m m m m m m m m d	04 04 04 04 04 04 04 04 04 04 04 04 04 0

SNP Base Columbia/ Landsberg	)			C/A	C/A															G/A	A/C	A/C	C/G	C/T	A/T	T/A	A/C	A/C	T/C	CT	A/T	T/A	G/A	A/T		
Indel Size Columbia/ Landsberg	TIL-	-2/2	1/-1			40/-40	3199/-3199	L/L-	36/-36	5451/-5451	-14/14	-4/4	14053/-	14053	-4/4	2388/-2388	-4/4	30/-30	6-/6																3/-3	-1/1
Method	2	-	1	1	1	2	2	7	7	2	2	2	2		2	2	2	2	2	1	1	<del></del> 1	1	1	<del></del>	_	1		<del></del>		1	1	1	1	2	1
Type	IND	ONI	IND	SNP	SNP	IND	IND	IND	ONI	ONI	IND	IND	IND		ON N	IND	ONI	IND	ONI	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	ONI	IND
Right	78012	4772	57542	75418	86109	12920	16333	19401	22629	28384	24107	26528	42196		47874	64076	62061	62751	82090	10776	10428	10467	10334	10362	10625	54112	71964	72088	72159	71701	34567	32758	17171	36226	7300	2075
Left	78011	4771	57540	75416	86107	12879	13133	19400	22592	22932	24106	26527	28142		47873	61687	62060	62720	82080	10774	10426	10465	10332	10360	10623	54110	71962	72086	72157	71699	34565	32756	17169	36224	7296	2074
Marker Name	469864	471206	471207	467921	468703	469865	469866	469867	469868	469869	469870	469871	469872		469873	469874	469875	469876	469877	471731	471732	471733	471734	471735	471736	472107	472164	472165	472166	472167	472170	472435	473109	473289	473959	474351
BAC Name	F20B17	F20B17	F20B17	F23H14	F23H14	F23H14	F23H14	F23H14	F23H14	F23H14	F23H14	F23H14	F23H14		F23H14	F23H14	F23H14	F23H14	F23H14	T5M2	T5M2	T5M2	T5M2	T5M2	T5M2	MJB20	MJB20	MJB20	MJB20	MJB20	MJB20	MJB20	MJB20	MJB20	MJB20	MJB20
BAC Chromosome Length		90149	90149	87584	87584	87584	87584	87584	87584	87584	87584	87584	87584		87584	87584	87584	87584	87584	94503	94503	94503	94503	94503	94503	82189	82189	82189	82189	82189	82189	82189	82189	82189	82189	82189
Chron	_	1		2	7	7	7	7	2	2	2	2	7		2	2	7	2	2	2	2	2	7	2	2	2	2	2	2	2	2	2	2	2	2	2
Seq id	AC010793	AC010793	AC010793	AC006837	AC006837	AC006837	AC006837	AC006837	AC006837	AC006837	AC006837	AC006837	AC006837		AC006837	AC006837	AC006837	AC006837	AC006837	AC007730	AC007730	AC007730	AC007730	AC007730	AC007730	AC007584	AC007584	AC007584	AC007584	AC007584	AC007584	AC007584	AC007584	AC007584	AC007584	AC007584
Seq	40	40	40	41	41	41	41	41	41	41	41	41	41		41	41	41	41	41	42	42	42	42	42	42	43	43	43	43	43	43	43	43	43	43	43

SNP Base Columbia/	Landsberg		A/C	G/C	T/C	T/G	C/T	T/C	A/G	A/T	G/T	A/G	A/G	T/A	G/C	J/L	D/L											T/A	A/G	G/A	C/A	S/C	A/G	C/T	$\mathrm{C/T}$	G/A	T/C
Indel Size Columbia/	Landsberg	1/-1																6-/6	-5/5	27/-27	4/-4	-4/4	3/-3	-14/14	5/-5	-4/4	-1/1										
	Method		1		_	1	1	_	_	_	_		<del></del>		_	1	1	2	2	2	2	2	2	2	2	2	_	-1	-	1	_	1	1	-	-		_
	Type		SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	ONI	ONI	ONI		ONI	ONI	ON.		ONI	ONI	SNP	SNP	SNP							
	Right	53757	7204	21548	25183	24284	24283	27216	26805	27004	27786	34752	57258	31602	31357	30475	36547	18799	19796	2323	23062	2641	2928	3325	53779	53790	35559	84817	85691	82282	81449	82927	80656	82828	81409	36818	37004
	Left	53755	7202	21546	25181	24282	24281	27214	26803	27002	27784	34750	57256	31600	31355	30473	36545	18789	19795	2295	23057	2640	2924	3324	53773	53789	32206	84815	82689	82280	81447	82925	80654	82826	81407	36816	37002
Marker	Name	474352	473036	472041	472294	472349	472350	472437	472438	472439	472440	472833	472874	472949	472950	472997	473167	474033	474034	474035	474036	474037	474038	474039	474040	474041	474476	472295	472296	472297	472298	472299	472300	472301	472302	472332	472333
BAC	Name	MJB20	T19E12	T7M7	T7M7	T7M7	T7M7	T7M7	T7M7	T7M7	T7M7	T7M7	T7M7	T7M7	T7M7	T7M7	T7M7	T7M7	T4P13	T4P13	T4P13																
BAC	Chromosome Length			57991	57991	57991	57991	57991	57991	57991	57991	57991	57991	57991	57991	57991	57991	57991	57991	57991	57991	57991	57991	57991	57991	57991	57991	93735	93735	93735	93735	93735	93735	93735			93735
	-		_	AC018721 2	AC018721 2	AC018721 2	AC018721 2	AC018721 2	AC018721 2	AC018721 2	AC018721 2	AC018721 2	AC018721 2	AC018721 2	AC018721 2	AC018721 2	AC018721 2	AC018721 2	AC008261 3	AC008261 3	AC008261 3																
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SNP Base Columbia/	Landsberg	T/A	T/C	C/G	T/G	A/G	C/T	C/T	G/T	A/T	T/C	C/G	C/T	A/T	A/G	G/A	C/A	T/C	G/C	T/C	G/A	A/G	C/T	G/C	A/T	G/A											
Indel Size Columbia/	Landsberg																										-4/4	1/-1	-1/1	-1/1	1/-1	1/-1	1/-1	-1/1	-2/2	-1/1	-1/1
	Method	-	1					_	-	1		_	1		1	-1					1	1	1	1	1	_	_	1	-	_	1	<b></b>	-	_		_	1
	Type	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	IND	ONI	IND	ONI	ONI						
	Right	57001	57542	74097	42784	42815	42699	78036	27226	30746	16551	16701	16538	16448	88268	20069	71122	68560	69752	71028	28971	28927	28991	66452	66983	35077	30775	32905	35559	42759	45923	65786	66187	66434	66562	70421	70997
	Left	56999	57540	77095	42782	42813	42697	78034	27224	30744	16549	16699	16536	16446	89386	69005	71120	68558	69750	71026	28969	28925	28989	66450	18699	35075	30774	32903	35558	42758	45921	65784	66185	66433	66561	70420	96601
Marker	Name	472338	472339	472362	472785	472786	472787	472816	472892	472935	473003	473004	473005	473006	473128	473464	473465	473466	473467	473468	473498	473499	473500	473503	473504	473701	474462	474463	474464	474465	474466	474467	474468	474469	474470	474471	474472
BAC	Name	T4P13	T4P13	T4P13	T4P13	T4P13	T4P13	T4P13	T4P13	T4P13	T4P13	T4P13	T4P13	T4P13	T4P13	T4P13	T4P13	T4P13	T4P13	T4P13	T4P13	T4P13	T4P13	T4P13	T4P13	T4P13	T4P13	T4P13	T4P13	T4P13	T4P13	T4P13	T4P13	T4P13	T4P13	T4P13	T4P13
BAC	Chromosome Length	93735	93735	93735	93735	93735	93735	93735	93735	93735	93735	93735	93735	93735	93735	93735	93735	93735	93735	93735	93735	93735	93735	93735	93735	93735	93735	93735	93735	93735	93735	93735	93735	93735	93735	93735	93735
			8261 3	8261 3	8261 3	8261 3	8261 3	8261 3	8261 3	18261 3	8261 3	8261 3	8261 3	8261 3	8261 3	8261 3	8261 3	8261 3	8261 3	_	_	8261 3		8261 3	8261 3	_	8261 3	_	_	8261 3	_	_	8261 3	8261 3	18261 3	18261 3	8261 3
_	n Seqid	AC008261	AC008261	AC00826	AC008261	AC00826	AC008261	AC00826	AC008261	AC008261	AC008261	AC008261	AC008261	AC00826	AC008261	AC00826	AC00826	AC008261	AC00826																		
Sed	unu	46	46	46	46	46	46	46	46	46	46	46	46	46	46	46	46	46	46	46	46	46	46	46	46	46	46	46	46	46	46	46	46	46	46	46	46

SNP Base Columbia/ Landsberg	C/A A/G	C/T A/T	A/G T/G	A/G A/T	T/A T/C	C/G A/G	T/G	A/C G/C	C/C	C/G T/A	T/C	A/G	G/A C/A	C/A	A/C	C/T	A/T	C/A	A/C	A/G	C/T	T/G	G/T
Indel Size Columbia/ Landsberg 2/-2 -1/1	-1/1																						
Method 1	<b>-</b>			<b></b>			· —	<del></del>	-		1	₩,		П	1	_		<b>—</b> ,	<b>.,</b>		1		<b>—</b>
Type IND IND	SNP SNP SNP	SNP	SNP	SNP SNP	SNP SNP	SNP	SNP	SNP SNP	SNP	SNP SNP	SNP	SNP	SNP SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP
Right 71087 82363	84002 10796 49032	77469 15175	15320 15320 15134	14557 16020	127045 126526	126534	127067	34116 32830	58169	19932 35725	36158	36310	17859 17259	17213	17405	17527	17401	93278	92943	28906	91584	1007	1736
Left 71084 82362	84001 10794 49030	77467 15173	15318 15318 15132	14555 16018	127043 126524	126532	127065	34114 32828	58167	19930 35723	36156	36308	17857 17257	17211	17403	17525	17399	93276	92941	58906	91582	1005	1734
Marker Name 474473	474475 466885 466963	466998 467131	467133 467133 467134	467135 467136	467181 467182	467183	467185	467367 467368	467442	467493 467512	467513	467514	467579 467580	467581	467582	467583	467584	467631	467632	467633	467634	467682	467683
BAC Name T4P13	T4P13 T12H1 T12H1	T12H1 T12H1	T12H1 T12H1 T12H1	T12H1 T12H1	T12H1 T12H1	T12H1 T12H1	T12H1	T12H1 T12H1	T12H1	T12H1 T12H1	T12H1	T12H1	T12H1 T12H1	T12H1	T12H1	T12H1	T12H1	T12H1	T12H1	T12H1	T12H1	T12H1	T12H1
BAC Chromosome Length 3 93735 3 93735	93735 140064 140064	140064	140064 140064 140064	140064 140064	140064 140064	140064	140064	140064	140064	140064 140064	140064	140064	140064 140064	140064	140064	140064	140064	140064	140064	140064	140064	140064	140064
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Seq id AC008261 AC008261	AC008261 AC009177 AC009177	AC009177 AC009177	AC009177 AC009177 AC009177	AC009177 AC009177	AC009177 AC009177	AC009177	AC009177	AC009177	AC009177	AC009177 AC009177	AC009177	AC009177	AC009177 AC009177	AC009177	AC009177	AC009177	AC009177	AC009177	AC009177	AC009177	AC009177	AC009177	AC009177
Seq num 46 46	46 74 74	74	4 4 4 7 4 7	47	47	74	4 4	47	47	47	47	41	47 74	47	47	47	47	47	47	47	47	47	47

SNP Base Columbia/ Landsberg A/C A/G G/T C/T T/A A/C G/A T/G A/G A/C	C/A
Indel Size Columbia/ Landsberg -3/3 13/-13 4/-4 242/-242 3/-3 -15/15 5/-5 3/-3 -1/1 -1/1 -2/2 -2/2 3/-3 -2/2 -2/2 -2/2 -2/2 3/-3	1/-1
Method  The state of the state	
	SNP
Right 2977 2537 86015 85652 67618 68903 9782 9783 65180 79433 107258 135453 12029 11410 136145 17843 4234 52288 52288 52335 77775 87758 96209 115173 16516 17262 17848 17853 91426	9787 85653
Left 2975 2535 86013 85650 67616 68901 9780 97771 11408 135451 107256 135451 11409 11409 11409 11409 115027 116027 136121 136121 136121 136121 136121 136121 136121 136121 136121 136121 136121 136121 17615 17777 16515 17777 16515	9785 85651
Marker Name 467684 467685 467837 467837 467838 467976 468311 468311 468312 468485 468411 468641 468936 470578 470581 470582 470583 471423 471423 471423	471695
BAC Name 1122H1	F22F7
	140064 91924
Chromosome	m m
Seq id AC009177	AC009606 AC009606
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SNP Base Columbia/	Landsberg	G/A	G/C	T/C	J/G	J/G	T/G	C/T	T/A	A/T	C/C	T/C	A/C	A/G	A/T	G/A	A/C	A/C	C/A	G/C	T/C	A/C	A/G	A/T	A/T	C/T	G/A	T/A	G/A	A/G	A/G	T/C	A/C	A/C	T/C	G/T	C/T
Indel Size Columbia/	Landsberg																																				
	Method	1	_		-	1	1	1	-		-1	-1	-		Ţ		1	1	1		<del></del>	-	-	-	1	-	_	<del></del>	1	1	1	1		1	-	1	1
	Type	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP						
	Right	85517	85720	85559	85737	85604	85533	85549	45473	45817	90312	89379	89779	90320	89801	52895	52826	31402	72856	72795	72750	72691	72877	72272	72866	54588	56725	56730	55493	56724	55901	39787	81393	5550	5036	5509	5448
	Left	85515	85718	85557	85735	85602	85531	85547	45471	45815	90310	89377	<i>LLL</i> 89777	90318	89799	52893	52824	31400	72854	72793	72748	72689	72875	72270	72864	54586	56723	56728	55491	56722	55899	39785	81391	5548	5034	5507	5446
Marker	Name	471696	471697	471698	471699	471700	471701	471702	472171	472172	472323	472324	472325	472326	472327	472466	472467	472482	472636	472637	472638	472639	472640	472641	472642	472733	472734	472735	472736	472737	472738	473152	473360	473368	473369	473370	473371
BAC	Name	F22F7	F22F7	F22F7	F22F7	F22F7	F22F7	F22F7	F22F7	F22F7	F22F7	F22F7	F22F7	F22F7	F22F7	F22F7	F22F7	F22F7	F22F7	F22F7	F22F7	F22F7	F22F7	F22F7	F22F7	F22F7	F22F7	F22F7	F22F7	F22F7	F22F7						
BAC	Chromosome Length	91924	91924	91924	91924	91924	91924	91924	91924	91924	91924	91924	91924	91924	91924	91924	91924	91924	91924	91924	91924	91924	91924	91924	91924	91924	91924	91924	91924	91924	91924	91924	91924	91924	91924	91924	91924
	Chro	n	33	3	$\kappa$	3	3	$\varepsilon$	3	$\epsilon$	3	3	33	$\epsilon$	æ	3	3	3	3	3	33	3	$\mathcal{C}$	3	3	33	33	$\varepsilon$	33	$\mathfrak{S}$	$\alpha$	3	3	$\epsilon$	33	3	3
	Seq id	AC009606	AC009606	AC009606	AC009606	AC009606	AC009606	AC009606	AC009606	AC009606	AC009606	AC009606	AC009606	AC009606	AC009606	AC009606	AC009606	AC009606	AC009606	AC009606	AC009606	AC009606	AC009606	AC009606	AC009606	AC009606	AC009606	AC009606	AC009606	AC009606	AC009606						
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SNP Base Columbia/ Landsberg A/T C/A T/G T/C		
Indel Size Columbia/ Landsberg	11/-11 6/-6 -3/3 -4/4 17/-17 -6/6 -4/4 -2/2 2/-2 1/-1 1/-1 1/-1 1/-1 1/-1 1/-1	-0/0 -12/12
Method 1 1 1 1 1	000000000000000000000000000000000000000	7 7
Type SNP SNP SNP SNP		N N
Right 5300 74553 11154 53949 38191	17389 27901 28945 32050 35059 3532 36310 37997 43376 46576 32052 32053 32053 72674 72770 10512 11767 13025 13236 37819 42042 58935 58935 58935 78137 78137	8719
Left 5298 74551 11152 53947 38189	17377 27894 28944 32049 35041 3531 36309 37931 46575 32051 32051 32052 32053 5262 7267 7267 10508 11763 13024 13235 37818 42038 58930 58930 58930 58930 58930 58930 58930	8718
Marker Name 473372 473515 473534 473657	473856 473858 473858 473860 473861 473862 473863 474110 474111 474111 474111 474111 474111 474111 470452 470453 470453 470454 470453 470454 470459 470460 470460	470464
BAC Name F22F7 F22F7 F22F7 F22F7	F22F7	MLP3
BAC Chromosome Length 3 91924 3 91924 3 91924 3 91924	91924 91924 91924 91924 91924 91924 91924 91924 91924 91924 91924 91924 91926 79296 79296 79296 79296	79296 79296
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Seq id AC009606 AC009606 AC009606 AC009606	AC009606 AC009176	AC009176 AC009176
Seq num 48 48 48 48	** ** * * * * * * * * * * * * * * * *	64

SNP Base Columbia/ Landsberg T/C A/G A/G G/A C/A A/G C/A A/G C/A A/G C/A A/G C/A A/G C/A A/G C/A A/C C/A C/A C/A C/A C/A C/A C/A C/A
Indel Size Columbia/ Landsberg  4/4 -9/9 7/-7 -4/4 -3/3 18/-18 -2/2 4/-4 -10/10 3/-3 6274/-6274 -4/4 -4/4 8/-8
Method 222222222222222222222222222222222222
Right 22735 23112 23482 233112 23482 23081 5443 5732 5733 5734 5737 34496 4702 26807 12204 14411 1448 112204 38246 30184 42097 42569 42097 47093 47093
Left 22733 23110 23480 23480 23079 5441 5730 5731 5735 34494 4700 28894 28298 26805 114410 14410 14410 14410 14709 37929 37929 37929 37929 42048 42048 43006 43046 43432 47092 47093 47093
Marker Name 466918 466919 466920 466921 468147 468148 468149 468149 468149 468149 468140 470402 470401 470404 470404 470405 470405 470404 470405 470406 471381 468110 468111 468110 468111 468111 468110 470473 470475 470475 470476 470477
BAC Name MCP4 MCP4 MCP4 MCP4 MCP4 MCP4 MCP4 MCP4
Chromosome Length  3
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Seq id AB028610 AB028617
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SNP Base Columbia/ Landsberg	C/T T/C A/C C/A T/A	G/C A/C 1/G	1/C 1/G 1/A 1/A A/T A/C
Indel Size Columbia/ Landsberg 8-8 7/-7 6/-6 -3/3 -3/3 -1/-1 -1/1	-44/44 -44/44 1221/-1221	8/-8 -13/13 -13/13 82/-82 -4/4 -1/1	
Method 2 2 2 2 1 1	000	00000	
Type CN CN C	ANS SANS ON		S S S S S S S S S S S S S S S S S S S
Right 48886 48989 49119 51110 5732 26252 41900	3178 1928 4381 7896 8667 10744 10753	13077 13636 13637 14988 9577 4283 14909 14963	15842 15809 20142 18233 9248 21188 21125 42935 7333
Left 48877 48981 49112 51109 51112 5722 26250 41899	3176 1926 4379 7894 8665 10743 10752	13068 13635 13636 14905 9576 4282 14924 14907 1303	15840 15807 20140 18231 9246 21123 42933 7331
Marker Name 470482 470483 470485 470485 470486 470487 471384	466800 467198 468112 468676 468677 470501 470503	470504 470505 470506 470507 471387 471679 471680 471681	471945 471946 472090 472424 472434 472835 473130 473259
BAC Name MOA2 MOA2 MOA2 MOA2 MOA2	MQD17 MQD17 MQD17 MQD17 MQD17 MQD17 MQD17	MQD17 MQD17 MQD17 MQD17 MQD17 K2019 K2019 K2019	K2019 K2019 K2019 K2019 K2019 K2019 K2019 K2019
BAC Chromosome Length  3	22199 22199 22199 22199 22199 22199 22199	22199 22199 22199 22199 22199 43500 43500 43500	43500 43500 43500 43500 43500 43500 43500 43500
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Seq num 51 51 51 51 51 51 51 51 51 51 51 51 51			23 23 23 23 23 23 23 23 23 23 23 23 23 2

SNP Base Columbia/ Landsberg A/C	C/G T/C C/T T/A A/C C/T	T/C T/A G/C A/G T/A T/A T/A
Indel Size Columbia/ Landsberg 1/-1 -1/1 -1/1 -4/4 8/-8 -5/5	-46/46 -78/78 -2/2 -1/1 -1/1 -1/1 2/-2	77-
Method 1 1 1 2 2	N — — — — — — — — — — — — — — — — — — —	
Type SNP IND IND SNP SNP IND		SNP SNP SNP SNP SNP SNP SNP
Right 29424 12418 42984 42985 42988 12287 12347	7765 42033 30928 23051 25996 24991 25839 24349 197 782 503 1011 17504 24170 24226 24236	38230 30684 66246 66082 47889 21817 21770 21770 21725
Left 29422 12416 42983 42984 42987 4661 12278	7764 42031 30926 23049 25837 24316 24347 195 780 502 1010 17503 24169 24225 24225	38229 30682 66244 66080 47887 21842 21768 21723 20915
Marker Name 473734 474314 474315 474316 474317 467314 470510	470512 471704 472116 472587 472705 472707 473539 473640 473640 473941 474320 474321 474322 474323	474525 471928 472065 472108 472124 472125 472127 472126
BAC Name K2019 K2019 K2019 K2019 K2019 K2019 K2019 K7012 MT012	MTO12 K24M9 K24M9 K24M9 K24M9 K24M9 K24M9 K24M9 K24M9 K24M9 K24M9 K24M9	K24M9 MMB 12 MMB 12 MMB 12 MMB 12 MMB 12 MMB 12 MMB 12
BAC Chromosome Length 3 43500 3 43500 3 43500 3 43500 3 19801 3 19801	19801 45292 45292 45292 45292 45292 45292 45292 45292 45292 45292 45292	45292 73977 73977 73977 73977 73977 73977
Chroi 3 3 3 3 3		
Seq id AB028608 AB028608 AB028608 AB028608 AB028620 AB028620	AB028620 AP001303	AP001303 AP000417 AP000417 AP000417 AP000417 AP000417 AP000417 AP000417
Seq 53 53 53 54 54	4	26 56 56 56 56 56 56 56 56 56 56 56 56 56

Indel Size SNP Base Columbia/ Columbia/	_	C/A	T/C	T/C	T/C	T/C	A/C	T/C	A/C	T/C	A/G	A/G	T/G	A/G	T/G	A/G	T/G	C/T	A/T	A/T	A/T	A/T	A/T	A/T	C/A	T/C	G/A	G/A	G/A	T/A	A/T	A/G	T/C	T/C	C/T	C/T	T/G
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	Right	25388	23434	23433	22955	22030	22000	21997	21865	21672	25896	22441	22032	21931	21923	21707	21688	25192	25077	22110	21883	21722	21714	25365	66069	68256	28856	28872	48275	14381	32813	6785	59992	62564	62200	40404	69562
	Left	25386	23432	23431	22953	22028	21998	21995	21863	21670	25894	22439	22030	21929	21921	21705	21686	25190	25075	22108	21881	21720	21712	25363	<i>L</i> 6069	68254	28854	28870	48273	14379	32811	6783	59990	62562	62198	40402	09569
Marker	Name	472129	472130	472131	472132	472133	472134	472135	472136	472137	472138	472139	472140	472141	472142	472143	472144	472145	472146	472147	472148	472149	472150	472151	472168	472169	472181	472182	472391	472425	472469	472477	472493	472507	472508	472564	472585
BAC	Name	MMB12	MMB12	<b>MMB12</b>	MMB12	MMB12	MMB12	MMB12	MMB12	<b>MMB12</b>	MMB12	MMB12	MMB12	<b>MMB12</b>	MMB12	MMB12	MMB12	<b>MMB12</b>	MMB12	MMB12	MMB12	MMB12	MMB12	MMB12	MMB12	<b>MMB12</b>	<b>MMB12</b>	<b>MMB12</b>	MMB12	MMB12	MMB12						
BAC	Chromosome Length	73977	73977	73977	73977	73977	73977	73977	73977	73977	73977	73977	73977	73977	73977	73977	73977	73977	73977	73977	73977	73977	73977	73977	73977	73977	73977	73977	73977	73977	73977	73977	73977	73977	73977	73977	73977
	Chro	3	$\epsilon$	33	3	3	3	3	3	3	3	3	3	3	3	3	3	3	3	3	3	3	3	3	3	3	$\mathcal{C}$	3	$\varepsilon$	3	3	3	В	3	$\epsilon$	3	33
	Seq id	AP000417	AP000417	AP000417	AP000417	AP000417	AP000417	AP000417	AP000417	AP000417	AP000417	AP000417	AP000417	AP000417	AP000417	AP000417	AP000417	AP000417	AP000417	AP000417	AP000417	AP000417	AP000417	AP000417	AP000417	AP000417	AP000417	AP000417	AP000417	AP000417	AP000417	AP000417	AP000417	AP000417	AP000417	AP000417	AP000417
Sed	umu	26	99	56	99	56	99	99	99	99	99	56	56	99	99	99	56	56	56	99	99	99	99	99	99	99	99	99	99	99	99	99	99	99	26	99	26

SNP Base	Londshara	C/G	T/G	G/T	G/T	C/A	G/A	T/G	G/A	G/C	G/T	C/T									A/C	C/G	CT	C/T	G/A	C/T	A/T	T/A	T/A	G/A	A/C	A/C	A/T	A/T	G/A	G/C	T/C
Indel Size	Londohara	Lamasocie											3/-3	3/-3	-3/3	-1/1	-4/4	-1/1	-1/1	-1/1																	
	Mathod	Incuror 1	1	1	-	-		_	1	1	<b>—</b>	1	2	2	2	<del></del>	1	1	_	1		_	_	<b>,</b>		_	_		_	_	1		_	_	_		-
	Ţ.	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	ONI	ONI	ONI	ONI	ONI	ONI	ONI	ONI	SNP																
	$D_{i, \alpha}^{c}$	70625	27618	27608	27632	50624	39000	39634	41060	13446	12412	12755	11079	16758	4724	22354	33542	7062	7063	71584	4318	5084	4252	5082	14934	14897	14912	35287	35259	34977	35255	34995	35286	35038	33874	34177	34344
	<del>4</del> ~ 1	70623	27616	27606	27630	50622	38998	39632	41058	13444	12410	12753	11075	16754	4723	22353	33541	7061	7062	71583	4316	5082	4250	5080	14932	14895	14910	35285	35257	34975	35253	34993	35284	35036	33872	34175	34342
Morbor	Momo	172586 472586	473033	473034	473035	473304	473651	473652	473793	473804	473805	473806	473962	473963	473964	474359	474360	474361	474362	474363	472061	472062	472063	472064	472290	472291	472292	472664	472665	472666	472667	472668	472669	472670	472671	472672	472673
ΣVΩ	Nome	MMB 12	MMB12	MMB12	MMB12	MMB12	MMB12	K10D20																													
C Y		3 73977	3 73977	3 73977	3 73977	3 73977	3 73977	3 73977	3 73977	3 73977	3 73977	3 73977	3 73977	3 73977	3 73977	3 73977	3 73977	3 73977	3 73977	3 73977	3 55161	3 55161			3 55161	3 55161	3 55161	3 55161	3 55161	3 55161	3 55161	3 55161	3 55161	3 55161	3 55161	3 55161	3 55161
	T: 0	Seq 10 AP000417	AP000417	AP000417	AP000417	AP000417	AP000417	AP000410																													
S	5	110111 56	56	56	99	26	99	99	56	56	99	26	99	99	26	99	26	99	99	99	57	57	27	57	57	27	57	57	27	57	57	57	57	57	57	57	27

SNP Base Columbia/ Landsberg A/G A/G T/A T/C C/T C/T C/G C/T T/A T/C C/T C/A	T/A G/A A/C A/C T/C
Indel Size Columbia/ Landsberg  Landsberg  1/-1 -1/1 -2/2 1/-1 1/-1	177
Method	
S S S S S S S S S S S S S S S S S S S	SNP SNP SNP SNP SNP
Right 34364 34519 111334 11140 111528 53958 54394 53903 40928 40683 16208 16591 16896 15354 15354 15356 39413 18595 37781 13801 27259 10598 10879 14918 14959	5005 16832 16708 5658 6231 6012
Left 34362 34517 11332 11138 11526 53956 54392 53901 40681 16206 16589 16589 16589 16589 16589 16589 16589 16589 16589 16589 16589 16589 1779 18593 13499 13469 115309 10596 10878 14917 14957	5008 16830 16706 5656 6229 6010
Marker Name 472674 472674 473208 473208 473209 473399 473510 473510 473531 473532 473531 473531 473531 473531 473531 473531 473531 473531 473531 473531 473531 473531 473531 473531 473531	4/4310 472039 472040 472072 472073
BAC Name K10020	K10D20 MHC9 MHC9 MHC9 MHC9
Chromosome Length  Chromosome Length  S5161  S5161	5 55161 3 58510 3 58510 3 58510 3 58510
	AP00410 AP001305 AP001305 AP001305 AP001305 AP001305
æ <b>8</b>	58 58 58 58 58 58

SNP Base Columbia/ Landsberg T/G T/G C/A A/T C/G G/T C/G G/T C/G G/T C/G C/A A/C A/C C/A A/C	1/G G/A A/T C/T 1/A
Indel Size Columbia/ Landsberg 4/-4 -4/4 -6/6 12/-12 -1/1 1/-1 3/-3 -4/4 1/-1 -1/1 -6/6 -4/4	
Method 1	
S S S S S S S S S S S S S S S S S S S	SNP SNP SNP SNP SNP SNP
Right 5889 37441 36725 36692 41451 31983 25642 25770 19103 50804 23041 23365 4416 542 21622 23364 36869 4428 48205	1981 22032 22033 12403 12668 15403
Left 5887 37439 36723 36690 41449 31981 25640 25768 19101 51497 51461 50429 50302 23032 23039 23592 2364 4415 529 2364 4415 529 2364 4415 4415 4415 4427 4427	1979 22030 22031 12401 12666 15401
Marker Name 472075 472184 472235 472235 472236 472923 472923 472973 472973 473965 473365 473365 473365 473365 473365 473365 473365 473365 473957 473957 473957 473958 474344 474345 474345 474345 474345 474345	468113 466874 466875 466934 466935
BAC Name MHC9 MHC9 MHC9 MHC9 MHC9 MHC9 MHC9 MHC9	MEK6 MZN24 MZN24 MZN24 MZN24 MZN24
BAC 58510	6184 82348 82348 82348 82348 82348
Chromosome 3 3 3 3 3 3 3 3 3 3 3 3 3 3 3 3 3 3 3	
Seq id AP001305	AP000739 AB028622 AB028622 AB028622 AB028622 AB028622
Seq. 258 258 258 258 258 258 258 258 258 258	60 60 60 60 60 60

SNP Base Columbia/	Landsberg	G/A	T/C	A/C	G/C	A/G	T/G	A/G	A/T	C/T	G/A	A/G	A/T	T/C	T/C	C/T	G/T	T/G	AT	C/T	T/G	G/A	A/C	A/T	C/T	C/T	G/A	A/C	A/G	CT	G/A						
Indel Size Columbia/	Landsberg																															-4/4	14/-14	-14/14	4/-4	12/-12	6/6-
	Method		<b>T</b>	_		_	_	1	_	-	_			_		-	-		1		_	<b>—</b>	1		1		<b>-</b>	1	_	-		2	2	2	2	2	2
	Type	SNP		ONI	ONI		ON!	ONI																													
	Right	14341	15224	15725	16086	15000	15402	14715	14960	13981	75592	75535	75466	21022	20839	62316	873	19463	19558	19530	33024	23244	23193	23078	23708	23853	46485	46601	46831	45797	55418	18076	25078	25583	41426	41465	41504
	Left	14339	15222	15723	16084	14998	15400	14713	14958	13979	75590	75533	75464	21020	20837	62314	871	19461	19556	19528	33022	23242	23191	23076	23706	23851	46483	46599	46829	45795	55416	18075	25063	25582	41421	41452	41503
Marker	Name	466937	466938	466939	466940	466941	466942	466943	466944	466945	467657	467658	467659	467706	467707	467812	468400	468415	468416	468417	468418	468419	468420	468421	468422	468423	468964	468965	468966	468967	469131	470521	470522	470523	470524	470525	470526
BAC	Name	MZN24																																			
BAC	Chromosome Length	3 82348	3 82348	3 82348	3 82348	3 82348	3 82348	3 82348	3 82348	3 82348	3 82348	3 82348	3 82348	3 82348	3 82348	3 82348	3 82348	3 82348	3 82348	3 82348	3 82348	3 82348	3 82348	3 82348	3 82348	3 82348	3 82348	3 82348	3 82348	3 82348	3 82348	3 82348	3 82348	3 82348	3 82348	3 82348	3 82348
	Seq id	AB028622																																			
Sea	_	09	09	09	9	09	09	99	09	09	09	09	09	9	09	09	09	09	09	09	09	09	09	09	09	09	09	09	9	09	09	09	09	9	09	09	09

SNP Base Columbia/ Landsberg	A/G A/T A/G C/G	7.4 T.7 A A T.7 A G.7 A G.
Indel Size Columbia/ Landsberg -9/9 713/-713 15609/- 15609 -9/9 -4/4 -7/7 2/-2 -1/1 -2/2 2/-2	77.	
Method 2 2 2 1 1 1 1		
	SNP SNP SNP SNP SNP SNP	SNP SNP SNP SNP SNP SNP SNP SNP SNP SNP
Right 41506 4908 57977 49983 5168 73431 15967 23850 28447 33232 46882 75605	887 40289 39456 8813 8924 8737	8737 19150 19432 5293 5357 5281 4942 29529 29490 29480 29605 29531 29531 17082 6171
Left 41505 4194 42367 49982 5167 73430 15964 23849 23849 2446 33229 46881 75603	880 40287 39454 8811 8922 8735	8735 19148 19430 5291 5355 5279 4940 29527 29478 29603 29529 29513 24555 16441 17080 6169
Marker Name 470527 470528 470529 470530 470531 471398 471399 471401 471402	471727 471727 471864 471869 471870 471871	471871 472191 472192 472207 472209 472209 472591 472594 472594 472595 472595 472596 473596 473357 473358
BAC	MZN24 MKA23 MKA23 MKA23 MKA23	MKA23
BAC Chromosome Length 3 82348 3 82348	82.348 70100 70100 70100 70100	70100 70100 70100 70100 70100 70100 70100 70100 70100 70100 70100
Chrom		
Seq id AB028622 AB028622 AB028622 AB028622 AB028622 AB028622 AB028622 AB028622 AB028622 AB028622 AB028622	AB028622 AP001306 AP001306 AP001306 AP001306	AP001306
Seq num 60 60 60 60 60 60 60 60 60 60 60	61 61 61 61 61	61 61 61 61 61 61 61 61 61 61 61 61 61 6

SNP Base Columbia/	Landsberg	A/T	T/C	A/G	A/G	C/G									T/C	A/G	G/A	T/G	A/T	G/A	C/T	T/A	T/G	T/C	T/A	T/A	A/C								G/A	CA	C/A
Indel Size Columbia/	Landsberg						750/-750	9/9-	1/-1	1/-1	-1/1	1/-1	-1/1	1/-1														3/-3	22/-22	4/-4	9/9-	-1/1	-1/1	-5/5			
	Method	_	1	1	<b>—</b>	-	2	2	_		-	_	1	-	_	-	-	1	-	-	1		1	<b>,</b> 1	<b>—</b>	_	_	2	2	2	2		_	_			1
	Type	SNP	SNP	SNP	SNP	SNP	ONI	ONI	ONI	IND	IND	ONI	ONI	ONI	SNP	SNP	SNP	SNP	ONI	ONI	ONI	IND	ONI	ONI	ONI	SNP	SNP	SNP									
	Right	6272	33097	33307	33607	60783	14779	529	1080	2700	39466	5300	5302	5314	2896	7045	38290	38726	38921	2703	43631	46628	46492	11637	45141	6044	6361	16172	463	8903	9450	26475	27910	30220	49539	49515	55822
	Left	6270	33095	33305	33605	60781	14028	528	1078	2698	39465	5298	5301	5312	5896	7043	38288	38724	38919	2701	43629	46626	46490	11635	45139	6042	6329	16168	440	8688	9449	26474	27909	30219	49537	49513	55820
Marker	Name	473443	473642	473643	473644	473673	473960	473961	474353	474354	474355	474356	474357	474358	471654	471852	472036	472037	472038	472102	472561	472776	472777	473115	473367	473677	473678	473838	473839	473840	473841	474072	474073	474074	471728	471729	472185
BAC	Name	MKA23	F16J14	F16J14	F16J14	F16J14	F16J14	F16J14	F16J14	F16J14	F16J14	F16J14	F16J14	F5N5	F5N5	F5N5																					
BAC	Chromosome Length	3 70100	3 70100	3 70100	3 70100	3 70100	3 70100	3 70100	3 70100	3 70100	3 70100	3 70100	3 70100	3 70100	3 47827	3 47827	3 47827	3 47827	3 47827	3 47827	3 47827	3 47827	3 47827	3 47827	3 47827	3 47827	3 47827	3 47827	3 47827	3 47827	3 47827	3 47827	3 47827	3 47827	3 71327	3 71327	3 71327
	Seq id (	AP001306	AP000731	AP000731	AP000731	AP000731	AP000731	AP000731	AP000731	AP000731	AP000731	AP000731	AP000731	AP001300	AP001300	AP001300																					
Seq	unu	61 ,	61	61 ,	61	61	61	61	61	61	61	61	61	61	62	7	62	62	62	62	62	7	7	62	62	62	7	, 29	62	62	62	. 62	62	. 62	63	63	. 63

SNP Base Columbia/	Landsberg	A/T	C/T	T/A	G/A	CT	A/C	T/A	T/G	G/T	T/G							T/C	G/T			A/T	T/A	A/C	A/C	T/A	T/A	CT	CT	A/T	G/T	AT	G/A	G/A	G/C	T/C	T/G
Indel Size Columbia/	Landsberg											1881/-1881	9/9-	8-/8	-1/1	-1/1	-1/1			-4/4	-4/4																
	Method	1			1	_	_	_	1	_	_	2	2	2	_		₩	-	-	2	2		-	Ţ			T	-	-	1	1			-			
	Type	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	IND	IND	ONI	IND	ONI	ONI	SNP	SNP	ONI	ONI	SNP															
	Right	55895	55397	32226	32235	29751	67202	19705	19704	25413	48239	14852	1472	21255	15958	24095	55627	1754	2138	458	459	61092	15248	15073	28666	27032	27031	33275	57376	76092	75839	63674	55689	55568	55673	55587	35411
	Left	55893	55395	32224	32233	29749	67200	19703	19702	25411	48237	12970	1471	21246	15957	24094	55626	1752	2136	457	458	61090	15246	15071	78664	27030	27029	33273	57374	26090	75837	63672	55687	55566	55671	55585	35409
Marker	Name	472186	472187	472307	472308	472661	473136	473295	473296	473427	473649	473920	473921	473922	474261	474262	474263	472887	472888	473879	473880	467302	467440	467441	467635	467687	467688	467799	468532	468581	468582	468600	468731	468732	468733	468734	468790
BAC	Name	F5N5	F5N5	F5N5	F5N5	F5N5	FSNS	F5N5	F5N5	F5N5	F5N5	F5N5	F5N5	F5N5	F5N5	F5N5	F5N5	F28F4	F28F4	F28F4	F28F4	MUJ8	MU18	MUJ8	MUJ8	MUJ8	MU18	MUJ8									
R AA	Chromosome Length	3 71327	3 71327	3 71327	3 71327	3 71327	3 71327	3 71327	3 71327	3 71327	3 71327	3 71327	3 71327	3 71327	3 71327	3 71327	3 71327	3 2490	3 2490	3 2490	3 2490	3 78921	3 78921	3 78921	3 78921	3 78921	3 78921	3 78921	3 78921	3 78921	3 78921	3 78921	3 78921	3 78921	3 78921	3 78921	3 78921
	Seq id (	300	AP001300	AP001300	AP001300	AP001300		AP001300	AP000733	AP000733	AP000733	AP000733	AB028621																								
Sec	T unu	63	63	63	63	63	63	63	63	63	63	63	63	63	63	63	63	64	49	49	49	65	65	65	65	65	65	65	65	65	65	65	65	65	65	65	65

SNP Base Columbia/	Landsberg	T/G	C/T	T/C	T/A	A/C	A/C	G/C	T/G	T/G	A/G	A/T	C/T	A/G																			G/C	T/C	A/T	G/A	T/C
Indel Size Columbia/	Landsberg														3/-3	9/9-	4/-4	-12/12	8-/8	4/-4	4/-4	-13/13	-1/1	1/-1	2/-2	-1/1	-1/1	-1/1	-1/1	1/-1	1/-1	-1/1					
	Method	1	_		<b>—</b>	<b></b>		-	1	_		-		_	2	2	2	2	2	2	2	2	_	1	_	<b></b>	1	1	_		_		_	_			Ţ
	Type	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	QNI	ONI	ONI		ONI		ONI	ONI	ONI	ONI	ONI	ONI	ON N	ONI	QN	ON.	ON ON	ONI	SNP	SNP	SNP	SNP	SNP
	Right	41781	42937	45707	70605	71795	70595	70603	72071	72085	90902	71796	70596	16880	15931	22566	39455	39507	48601	80902	71133	75230	1197	15931	15935	23748	25687	70597	2000	20907	20698	73275	34701	38382	38381	39779	39047
	Left	41779	42935	45705	70603	71793	70593	70601	72069	72083	70604	71794	70594	16878	15927	22565	39450	39506	48592	70603	71128	75229	1196	15929	15932	23747	25686	70596	70599	90902	96902	73274	34699	38380	38379	39777	39045
Marker	Name	469082	469083	469107	469183	469184	469185	469186	469187	469188	469189	469190	469191	469323	470513	470514	470515	470516	470517	470518	470519	470520	471388	471389	471390	471391	471392	471393	471394	471395	471396	471397	466917	467637	467638	467639	467640
BAC	Name	MUJ8	MUJ8	MUJ8	MUJ8	MUJ8	MUJ8	MUJ8	MUJ8	MUJ8	MUJ8	MUJ8	MUJ8	MUJ8	MUJ8	MUJ8	MUJ8	MUJ8	MUJ8	MUJ8	MUJ8	MUJ8	MUJ8	MUJ8	MUJ8	MUJ8	MUJ8	MUJ8	MUJ8	MUJ8	MUJ8	MUJ8	MSD24	MSD24	MSD24	MSD24	MSD24
BAC	Chromosome Length	78921	78921	78921	78921	78921	78921	78921	78921	78921	78921	78921	78921	78921	78921	78921	78921	78921	78921	78921	78921	78921	78921	78921	78921	78921	78921	78921	78921	78921	78921	78921	40018	40018	40018	40018	40018
	כ	3	ĸ	33	$\alpha$	$\varepsilon$	$\boldsymbol{\omega}$	$\varepsilon$	$\kappa$	E	æ	$\mathcal{C}$	33	3	$\omega$	æ	$\varepsilon$	æ	$\epsilon$	$\kappa$	3	E	ς.	3	33	33	33	33	€.	8	33	3		'n			<u>ω</u>
	Seq id	AB028621	AB028621	AB028621	AB028621	AB028621	AB028621	AB028621	AB028621	AB028621	AB028621	AB028621	AB028621	AB028621	AB028621	AB028621	AB028621	AB028621	AB028621	AB028621	AB028621	AB028621	AB028621	AB028621	AB028621	AB028621	AB028621	AB028621	AB028621	AB028621	AB028621	AB028621	AP000740	AP000740	AP000740	AP000740	AP000740
Sed	umu	65	65	65	65	65	65	65	65	65	65	65	65	65	65	65	65	65	65	65	65	65	65	65	65	65	65	65	. 59	65	65	65	99	99	99	99	99

SNP Base Columbia/	Landsberg	A/G	T/G	T/C		T/C	A/G	T/C	C/T	G/T	T/A	A/G	C/A	G/A	T/C	C/T	C/T	G/C	G/A	T/C	A/G	A/T	G/A	A/C	A/T	CT	T/A	C/T	A/T	G/A	G/A	T/C	A/G	A/G	A/T	G/C	C/A
Indel Size Columbia/	Landsberg				11/-11																																
	Method		_	_	2	П	-	_	_	-	-	1		_		_	-			П		1	-		_	<b>—</b>	-	-	-	-		1		_			
	Type	SNP	SNP	SNP	Q N	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP													
	Right	39759	15512	35222	22206	52760	52845	46999	46661	46643	46155	45791	63372	63442	63402	62844	63444	44803	64920	65570	65584	64925	58820	58851	58902	58357	50020	50134	50112	47778	47727	47733	47705	47736	47682	5277	19649
	Left	39757	15510	35220	22194	52758	52843	46997	46659	46641	46153	45789	63370	63440	63400	62842	63442	44801	64918	65568	65582	64923	58818	58849	58900	58355	50018	50132	50110	47776	47725	47731	47703	47734	47680	5275	19647
Marker	Name	467641	467840	468363	470509	471648	471649	471741	471742	471743	471783	471784	471787	471788	471789	471790	471791	471818	471848	471849	471850	471851	471857	471858	471859	471860	471888	471889	471890	471992	471993	471994	471995	471996	471997	472347	472357
BAC	Name	MSD24	MSD24	MSD24	MSD24	K7P8	K7P8	K7P8	K7P8	K7P8	K7P8	K7P8	K7P8	K7P8	K7P8	K7P8	K7P8	K7P8	K7P8	K7P8	K7P8	K7P8	K7P8	K7P8													
BAC	Chromosome Length	3 40018	3 40018	3 40018	3 40018	3 78535	3 78535	3 78535	3 78535	3 78535	3 78535	3 78535	3 78535	3 78535	3 78535	3 78535	3 78535	3 78535	3 78535	3 78535	3 78535	3 78535	3 78535	3 78535	3 78535	3 78535	3 78535	3 78535	3 78535	3 78535	3 78535	3 78535	3 78535	3 78535	3 78535	3 78535	3 78535
	Seq id	AP000740	AP000740	AP000740	AP000740	AB028609	AB028609	AB028609	AB028609	AB028609	AB028609	AB028609	AB028609	AB028609	AB028609	AB028609	AB028609	AB028609	AB028609	AB028609	AB028609	AB028609	AB028609	AB028609													
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SNP Base Columbia/	Landsberg	C/A	T/A	G/C	A/G	G/A	C/A	C/G	C/G	ΑΛΤ	C/A	T/C	A/C	A/C	T/C	A/C	C/G	A/G	C/T	T/C	CT	T/A	A/G	AVT	T/C	G/C	T/G	C/A	C/A	T/C	A/G	J/G	C/T	T/A	G/C	T/A	T/A
Indel Size Columbia/	Landsberg																																				
	Method		1	1	-	_		-	<b>-</b>	1	-	_	_	-		1	-		1	<b></b>			1	1	<b>—</b> 4	т				_		1		_			<b>~</b>
	Type	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP
	Right	19650	19858	20069	19926	32588	32709	32406	32407	32393	11609	11207	11315	11351	11356	11403	9964	10290	10531	39421	38765	23789	23790	23791	30592	30478	30306	1822	2098	1981	1901	1697	29140	69895	56947	7438	7451
	Left	19648	19856	20067	19924	32586	32707	32404	32405	32391	11607	11205	11313	11349	11354	11401	3966	10288	10529	39419	38763	23787	23788	23789	30590	30476	30304	1820	2096	1979	1899	1695	29138	26867	56945	7436	7449
Marker	Name	472358	472359	472360	472361	472408	472409	472410	472411	472412	472456	472457	472458	472459	472460	472461	472462	472463	472464	472766	472767	472828	472829	472830	473040	473041	473042	473063	473064	473065	473066	473067	473137	473277	473278	473516	473517
BAC	Name	K7P8	K7P8	K7P8	K7P8	K7P8	K7P8	K7P8	K7P8	K7P8	K7P8	K7P8	K7P8	K7P8	K7P8	K7P8	K7P8	K7P8	K7P8	K7P8	K7P8	K7P8	K7P8	K7P8	K7P8	K7P8	K7P8	K7P8	K7P8	K7P8	K7P8	K7P8	K7P8	K7P8	K7P8	K7P8	K7P8
BAC	Chromosome Length	78535	78535	78535	78535	78535	78535	78535	78535	78535	78535	78535	78535	78535	78535	78535	78535	78535	78535	78535	78535	78535	78535	78535	78535		78535		78535	78535	78535	78535	78535	78535	78535	78535	78535
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SNP Base Columbia/ Landsberg T/A T/C T/C A/C C/T	1/C C/T 1/C G/T G/T T/A G/C
Indel Size Columbia/ Landsberg 7/-7 3/-3 -20/20 -4/4 -4/4 6/-6 -3/3 4/-4 3/-3 -1/1 1/-1 1/-1 1/-1 -2/2 -1/1 -2/2 -1/1 -2/2 -1/1 -2/2 -1/1 -1/1	1-/1
Method	
	SNP SNP SNP SNP SNP SNP SNP SNP
Right 5914 5785 5796 6445 7247 6449 11497 15251 17674 17785 18956 24577 4408 11498 20206 30384 32398 32403 38752 4423 6198	63106 41583 35575 59692 59708 59516 47771 72456 72690
Left 5912 5783 5783 5794 6443 7245 6447 11489 11784 11784 11895 24570 4407 11499 20205 30383 32396 32401 38751 44747 59400 59400	63104 41581 35573 59690 59706 59514 47769 72454 72688
Marker Name 473518 473520 473521 473521 473523 473942 473943 473944 473945 473946 473946 473946 473946 474320 474320 474331 474333 474333 474333 474333	474339 471687 471807 472380 472381 472407 472629 472630
BAC Name K7P8 K7P8 K7P8 K7P8 K7P8 K7P8 K7P8 K7P8	K7P8 K13N2 K13N2 K13N2 K13N2 K13N2 K13N2 K13N2 K13N2 K13N2
BAC Chromosome Length 3 78535 3 7855 7855 7855 7855 7855 7855 7855 7855 7855 7855 7855 7855 7855 7855 7855 7855 785 78	78535 77483 77483 77483 77483 77483 77483
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Seq id AB028609	AB028609 AB028607 AB028607 AB028607 AB028607 AB028607 AB028607 AB028607 AB028607
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SNP Base Columbia/	Landsberg	A/T	C/T	C/T	T/A	A/G	C/T					A/T	CŢ	C/I	G/A	T/C	G/A	C/G	A/G	A/T	S/O	C/G	G/A	C/G	T/G	T/C	A/T	G/A	G/A	C/G	T/C	A/C	T/A	A/G	T/C	G/C	C/C
Indel Size Columbia/	Landsberg							9/9-	-3/3	16/-16	-1/1																										
	Method		-	1	_		_	2	2	2	1			1	_	1	-	-	_	-	-	-	1			₩.	<b>—</b>	1	<b>-</b>	1	1	1	*****		_	_	<b>-</b>
	Type	SNP	SNP	SNP	SNP	SNP	SNP	ONI	QNI	ONI	ON.	SNP																									
	Right	72444	72543	72674	28930	28947	28984	10371	12417	14544	35556	4655	5149	4948	66985	58395	13443	13395	13379	13562	20706	68473	54117	53843	54113	30262	30323	43598	43520	62301	70736	27358	19775	20047	52764	52759	52577
	Left	72442	72541	72672	28928	28945	28982	10370	12416	14527	35555	4653	5147	4946	28697	58393	13441	13393	13377	13560	50704	68471	54115	53841	54111	30260	30321	43596	43518	62299	70734	27356	19773	20045	52762	52757	52575
Marker	Name	472632	472633	472634	472801	472802	472803	473932	473933	473934	474311	471703	471777	471778	471926	471927	472019	472020	472021	472022	472572	472757	472812	472813	472814	472831	472832	472905	472906	472982	473068	473108	473490	473491	473629	473630	473631
BAC	Name	K13N2	F20C19																																		
BAC	Chromosome Length	3 77483	3 77483	3 77483	3 77483	3 77483	3 77483	3 77483	3 77483	3 77483	3 77483	3 71184	3 71184	3 71184	3 71184	3 71184	3 71184		3 71184	3 71184	3 71184	3 71184	3 71184	3 71184	3 71184	3 71184	3 71184	3 71184	3 71184	3 71184	3 71184	3 71184	3 71184	3 71184	3 71184	3 71184	3 71184
	Seq id	AB028607	AP001298																																		
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SNP Base Columbia/	Landsberg	1/C	A/C	A/G	CT	A/T					A/G	C/T	C/T	T/A	G/A	C/T	T/A	A/G	C/T	T/C	A/T	G/A	CT	G/A													
Indel Size Columbia/	Landsberg						38/-38	-1/1	-1/1	1/-1																10/-10	-3/3	12277/- 12277	29/-29	29/-29	1-12	1193/-1193	117.11	11/-11	-10/10	11/-11	-3/3
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	Right	52446	52405	52712	52692	52682	6211	16828	41067	50771	80357	80297	79920	15120	15557	15060	65828	65613	65715	75234	75507	3958	3893	39563	39818	10103	14366	28746	27161	27162	28830	33179	360	300	36050	379	4454
	Left	52444	52403	52710	52690	52680	6172	16827	41066	50769	80355	80295	79918	15118	15555	15058	65826	65611	65713	75232	75505	3956	3891	39561	39816	10092	14365	16468	27131	27132	28822	31985	240	348	36049	367	4453
Marker	Name	473632	473633	473634	473635	473636	473854	474104	474105	474106	466957	466958	466959	466976	466977	466978	467322	467323	467324	468085	468086	468778	468779	469262	469263	470407	470408	470409	470410	470411	470412	470413	470414	4/0414	470415	470416	470417
BAC	Name	F20C19	MFE16	MFE16	MFE16	<b>MFE16</b>	MFE16	MFE16	MFE16	MFE16	MFE16	MFE16	MFE16	<b>MFE16</b>	MFE16	MFE16	MFE16	MFE16	MFE16	MFE16	MEE16	MFE16	MFF16	MFF16	MEET	MFE16	MFE16	MFE16	MFE16								
BAC	hromosome	71184		71184	71184	71184	71184	71184	71184	71184	82646	82646	82646	82646	82646	82646	82646	82646	82646	82646	82646	82646	82646	82646	82646	82646	3 82646	3 82646	97908						3 82646		3 82646
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SNP Base Columbia/ Landsberg T/A T/A T/A T/C C/G C/A T/C A/G A/G	A/T A/C A/G A/T A/G C/G
Indel Size Columbia/ Landsberg 7/-7 9/-9 8/-8 16/-16 16/-16 16/-16 5/-5 -15/15 -15/15 12/-12 -9/9 1/-1 1/-1 1/-1 1/-1 2/-9/9 19/-19 12/-12 -6/6 4/-4 8/-8	
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	SNP SNP SNP SNP SNP SNP SNP
Right 61638 63523 71764 744 745 74052 77825 77825 77825 75267 75267 75335 38283 3828	78266 60216 60330 21476 21279 64849 64278
Left 61630 63513 71755 727 727 728 74046 77824 78953 79710 75265 75333 38248 66535 27368 16568 31872 47865 56619 68665 73388	78264 60214 60328 21474 21277 64847 64276 64219
Marker Name 470418 470420 470421 470422 470423 470423 470425 470425 470425 470427 471382 470427 471383 467399 467399 46813 468135 470465 470465 470465 470465 470465 470465	467605 468067 468072 468073 468784 468785 468785
BAC Name MFE16 MFE	MODI MODI MODI MODI MODI MODI MODI
BAC Chromosome Length 3 82646 3 86139 3 86139 3 86139 3 86139 3 86139 3 86139 3 86139 3 86139 3 86139 3 86139	
Seq id AB028611 AB028611 AB028611 AB028611 AB028611 AB028611 AB028611 AB028616	AB028618 AB028618 AB028618 AB028618 AB028618 AB028618 AB028618
Seq numm 70 70 70 70 70 70 70 70 70 70 70 70 70	57 57 57 57 57 57 57 57 57 57 57 57 57 5

SNP Base Columbia/ Landsberg	17.G C/G G/C G/A A/C G/T C/A A/C C/A A/C C/G C/G T/A T/G A/T	
Indel Size Columbia/ Landsberg -7/7 4/-4 -10/10 106/-106 -10/10 162/-162 121/-121 4/-4 52/-52 52/-52 52/-52	7410-11410	75/-75 75/-75 -23/23 -1/1
Method 2 2 2 2 2 2 2 2 2 2 2 2 2 2 2 2 2 2 2	,	7 2 2 7
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Right 12859 1452 34069 36175 43346 52023 66853 73646 80872 82811 84749	72688 72664 76864 1270 7691 7749 34967 34967 34958 35098 207 75565 75705 38275 38275 38275 48310	2393 2393 9599 48314
Left 12858 1447 34068 36068 43345 51860 66731 73641 80819 82758 84696	7459 72662 76862 1268 7689 7747 34965 34965 34965 35096 205 75703 38273 38273 38273 48716 48308	2314 2317 9598 48313
Marker Name 470488 470489 470490 470491 470493 470494 470496 470499	471646 471647 471745 471745 471862 471952 471953 471955 471956 472017 472278 472278 472278 472278 472278 472278	473935 473936 473937 474312
BAC Name MODI MODI MODI MODI MODI MODI MODI MODI		K17E7 K17E7 K17E7 K17E7
BAC Chromosome Length 3 85690 3 85690 3 85690 3 85690 3 85690 3 85690 3 85690 3 85690 3 85690 3 85690	\$2090 \$3 \$2356 \$3 \$3 \$2356 \$3 \$2356 \$3 \$2356 \$3 \$2356 \$3 \$2356 \$3 \$2356 \$3 \$2356 \$3 \$2356 \$3 \$2356 \$3 \$2356 \$3 \$2356 \$3 \$2356 \$3 \$2356 \$3 \$25 \$25 \$25 \$25 \$25 \$25 \$25 \$25	3 82356 3 82356 3 82356 3 82356
Seq id AB028618		AP000736 AP000736 AP000736 AP000736
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SNP Base Columbia/ Landsberg A/C G/C	C/A G/A T/G A/G C/T C/T	1/C 1/C C/G G/T C/T
Indel Size Columbia/ Landsberg 1/-1 3/-3 8/-8 1142/-1142 10/-10 7/-7 8/-8 12/-12 -4/4	3/-3 -4/4 5/-5 4/-4 12/-12 -1/1 2/-2 2/-2 865/-865	12/-12
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Type SNP SNP SNP SNP SNP SNP SNP SNP SNP SNP	GNI GNI GNI GNI GNI GNI GNI GNI GNI GNI	SNP SNP SNP SNP SNP SNP SNP
Right 48374 17325 17021 10022 13011 63377 6502 65805 65912	77854 77985 81960 82102 5118 5495 5745 5657 5676 5472 5619 379 4931 5613	78180 81170 81116 81119 80708 81168 80699 15135
Left 48372 17323 17019 10218 13002 62234 6308 6494 65796 65859	77850 77850 77984 81954 82097 5116 5493 5274 5470 5617 366 4930 5510	78167 81168 81114 81117 80706 81166 80698 15133
Marker Name 474313 467375 467376 470440 470441 470442 470444 470445	470448 470449 470449 470451 471770 471771 471773 471774 471775 471775 471775 471776 471776 474109	470438 472211 472212 472213 472214 472215 474407
BAC Name K17E7 MIL15	MIL.15 MIL.15 MIL.15 MIL.15 MIL.15 F21A17	MIF6 T18B22 T18B22 T18B22 T18B22 T18B22 T18B22
BAC Chromosome Length 3 82356 3 84157 3 84157 3 84157 3 84157 3 84157 3 84157 3 84157 3 84157 3 84157	84157 84157 84157 84157 84157 64714 64714 64714 64714 64714 64714 64714 64714 64714 64714	82347 91567 91567 91567 91567 91567 91567
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Seq id AP000736 AB028615 AB028615 AB028615 AB028615 AB028615 AB028615 AB028615 AB028615	AB028615 AB028615 AB028615 AB028615 AP000732	AB028614 AL138652 AL138652 AL138652 AL138652 AL138652 AL138652
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SNP Base Columbia/	Landsberg	I/G	C/G	C/I	G/A	A/C	A/T	G/A	A/T	G/A	G/C	G/C	A/G	C/T	G/A	G/A	T/C	C/T	C/T	A/G	A/G	A/G	T/C	G/A	C/C	A/G	A/G	G/C	A/T	A/T	T/A	A/C	G/T	G/A	A/G	T/A	T/A
Indel Size Columbia/	Landsberg																																				
	Method	<b>—</b>		1	-	-	1	, <b>.</b>		-	1	-1				1	1		1	1	1	-		<b></b>		_		1	-	_		_	1	_	_	_	
	Type	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP						
	Right	15089	14982	44678	28615	28955	28938	69971	70225	64463	64451	64304	64343	64477	85095	85274	85096	84955	40022	42498	42499	41980	60237	49970	50018	49837	99009	81375	81297	81299	61933	61898	61876	80332	79733	12149	12159
	Left	15087	14980	44676	28613	28953	28936	69669	70223	64461	64449	64302	64341	64475	85093	85272	85094	84953	40020	42496	42497	41978	60235	49968	50016	49835	50064	81373	81295	81297	61931	61896	61874	80330	79731	12147	12157
Marker	Name	471828	471829	472160	472304	472305	472306	472378	472379	472392	472393	472394	472395	472396	472489	472490	472491	472492	472676	472889	472890	472891	472926	472992	472993	472994	472995	473493	473494	473495	473535	473536	473537	473542	473543	473681	473682
BAC	Name	T14D3	T14D3	T14D3	T14D3	T14D3	T14D3	T14D3	T14D3	T14D3	T14D3	T14D3	T14D3	T14D3	T14D3	T14D3	T14D3	T14D3	T14D3	T14D3	T14D3	T14D3	T14D3	T14D3	T14D3	T14D3	T14D3		T14D3	T14D3	T14D3						
BAC	hromosome	3 88010	3 88010	3 88010	3 88010	3 88010	3 88010	3 88010	3 88010	3 88010	3 88010	3 88010	3 88010	3 88010	3 88010	3 88010	3 88010	3 88010	3 88010	3 88010	3 88010	3 88010	3 88010	3 88010	3 88010	3 88010	3 88010	3 88010	3 88010	3 88010	3 88010	3 88010	3 88010	3 88010	3 88010	3 88010	3 88010
		AL138649	AL138649	AL138649	AL138649	AL138649	AL138649	AL138649	AL138649	AL138649	AL138649	AL138649	AL138649	AL138649	AL138649	AL138649	AL138649	AL138649	AL138649	AL138649	AL138649	AL138649	AL138649	AL138649	AL138649	AL138649	AL138649	AL138649	AL138649	AL138649	AL138649						
Seq	unu	78	78	78	78	78	78	78	78	78	78	78	78	78	78	78	78	78	78	78	78	78	78	78	78	78	78	78	78	78	78	78	78	78	78	78	78

SNP Base Columbia/	Landsberg	T/A	A/T	A/I	A/T	C/G	T/G								G/A	G/A	G/A	T/C	C/T	T/A	G/A	A/G	CT	CT	T/G	G/A	A/I										
Indel Size Columbia/	Landsberg							1/-1	-1/1	-1/1	1/-1	1/-1	-1/1	1/-1														-3/3	-11/11	9/9-	LIL-	L/L-	13/-13	9-/9	9/9-	-10/10	-10/10
	Method		1			1	1	1	1	_	1	_		1				-	-	-	<b>-</b>	<b>.</b>				_		2	2	2	2	2	2	7	7	5	2
	Type	SNP	SNP	SNP	SNP	SNP	SNP	IND	ON N	ONI	IND	ON N		IND	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	ONI	ONI	ON.	ONI	ONI	ONI	ONI			ON N
	Right	12160	12153	12154	12155	30967	30504	12156	28846	30948	49832	50138	81377	84997	49851	926	1155	217	836	34459	34319	34494	34426	35063	33073	38978	39206	13016	13161	19559	25308	25319	46600	47286	47339	47390	48397
	Left	12158	12151	12152	12153	30965	30502	12154	28845	30947	49830	50136	81376	84995	49849	974	1153	975	834	34457	34317	34492	34424	35061	33071	38976	39204	13015	13160	19558	25307	25318	46586	47279	47338	47389	48396
Marker	Name	473683	473684	473685	473686	473715	473716	474375	474376	474377	474378	474379	474380	474381	467042	467388	467389	467390	467391	468103	468104	468105	468106	468107	468334	469279	469280	469768	469769	469770	469771	469772	469773	469774	469775	469776	469777
BAC	Name	T14D3	T14D3	T14D3	T14D3	T14D3	T14D3	T14D3	T14D3	T14D3	T14D3	T14D3	T14D3	T14D3	F18N11	F18N11	F18N11	F18N11	F18N11	F18N11	F18N11	F18N11	F18N11	F18N11	F18N11	F18N11	F18N11	F18N11	F18N11	F18N11	F18N11	F18N11	F18N11	F18N11	F18N11	F18N11	F18N11
BAC	Chromosome Length	3 88010	3 88010	3 88010	3 88010	3 88010	3 88010	3 88010	3 88010	3 88010	3 88010	3 88010	3 88010	3 88010	3 91274	3 91274	3 91274	3 91274	3 91274	3 91274	3 91274	3 91274	3 91274	3 91274	3 91274	3 91274	3 91274	3 91274	3 91274	3 91274	3 91274	3 91274	3 91274	3 91274	3 91274		3 91274
	Sea id	649	AL138649	AL132953	AL132953	AL132953		AL132953	AL132953	AL132953	AL132953		AL132953																								
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SNP Base Columbia/	Landsberg													G/A	T/A	G/A	G/A	C/A	T/C	T/C	A/C	A/C	A/G	A/G	G/A	G/C	A/G	C/T		T/A	G/C	A/C	A/G	A/G			
Indel Size Columbia/	Landsberg	-3/3	-3/3	23/-23	-15/15	4/-4	L/L-	6-/6	-4/4	-4/4	3/-3	-1/1	1/-1																-1/1						-5/5	404/-404	3/-3
	Method	2	2	2	2	2	2	2	2	2	2	1			_	_	1		1	_			_	_	-		_	<b>—</b>	1	П		1	1	1	2	2	2
	Type		ONI	ON.		ONI	IND	ONI	ONI	ONI	ONI	ONI	IND	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	ONI	SNP	SNP	SNP	SNP	SNP	QNI	ON.	ONI
	Right	48525	49102	72221	75761	76045	78098	84186	89931	89932	90236	33022	878	3731	3696	3687	3663	3636	3768	3707	3690	3681	3648	3909	18207	18467	74942	75086	18389	79443	79559	79615	79455	79511	11609	73600	97564
	Left	48524	49101	72197	75760	76040	78097	84176	89930	89931	90232	33021	876	3729	3694	3685	3661	3634	3766	3705	3688	3679	3646	3907	18205	18465	74940	75084	18388	79441	79557	79613	79453	79509	11608	73195	97560
Marker	Name	469778	469779	469780	469781	469782	469783	469784	469785	469786	469787	471186	471187	471877	471878	471879	471880	471881	471882	471883	471884	471885	471886	471887	472094	472095	473798	473799	474301	468139	468140	468141	468142	468143	469761	469762	469763
BAC	Name	F18N11	F18N11	F18N11	F18N11	F18N11	F18N11	F18N11	F18N11	F18N11	F18N11	F18N11	F18N11	F9K21	F9K21	F9K21	F9K21	F9K21	F9K21	F9K21	F9K21	F9K21	F9K21	F9K21	F9K21	F9K21	F9K21	F9K21	F9K21	F18L15	F18L15	F18L15	F18L15	F18L15	F18L15	F18L15	F18L15
BAC	Chromosome Length	91274	91274	91274	91274	91274	91274	91274	91274	91274	91274	91274	91274	110980	110980	110980	110980	110980	110980	110980	110980	110980	110980	110980	110980	110980	110980	110980	110980	100328	100328	100328	100328	100328	100328	100328	100328
	Chr	$\mathcal{C}$	$\epsilon$	3	$\epsilon$	$\epsilon$	33	3	$\epsilon$	$\epsilon$	$\epsilon$	3	$\epsilon$	3	3	3	$\mathcal{E}$	$\varepsilon$	$\epsilon$	$\varepsilon$	$\epsilon$	$\varepsilon$	$\varepsilon$	$\varepsilon$	$\varepsilon$	$\varepsilon$	$\epsilon$	3	$\epsilon$	n	33	3	3	$\epsilon$	3	33	B
	Seq id	AL132953	AL132953	AL132953	AL132953	AL132953	AL132953	AL132953	AL132953	AL132953	AL132953	AL132953	AL132953	AL138657	AL138657	AL138657	AL138657	AL138657	AL138657	AL138657	AL138657	AL138657	AL138657	AL138657	AL138657	AL138657	AL138657	AL138657	AL138657	AL133298	AL133298	AL133298	AL133298	AL133298	AL133298	AL133298	AL133298
Seq	unu	79	79	79	79	79	79	79	6/	79	79	79	79	80	80	80	80	80	80	80	80	80	80	80	80	80	80	80	80	81	81	81	81	81	81	81	81

SNP Base Columbia/ Landsberg	G/A C/A G/A 17/A G/A G/T	G/T 1/A 1/A 1/A 1/A 1/A 1/A 1/A 1/A 1/A
Indel Size Columbia/ Landsberg 30/-30 305/-305 -4/4 -5/5	·	3/-3 305/-305 -4/4 -5/5 -21/21 -21/21 12/-12 -4/4 3/-3 807/-807
Method 2 2 2 2 2		
Type GNI GNI GNI GNI CNI	SNP SNP SNP SNP SNP SNP SNP	
Right 97855 98553 99105 99150 88384	36394 62001 62150 62253 60591 62257	62029 62262 65708 65478 65486 25669 25693 91003 26407 26408 37558 37550 38420 59417 10480 11469 12021 12021 13232 13234 13117 21556
Left 97824 98247 99104 99149	36392 61999 62148 62251 60589 62255	62027 62260 65706 65484 25674 25691 25691 26405 37548 38418 38418 59415 10476 11163 12020 12065 13231 13233 13704 21555
Marker Name 469764 469765 469766 469767	467656 468057 468058 468060 468061 468061	468062 468063 468216 468218 468298 468293 469145 469146 469309 469309 469476 469309 469542 469543 469543 469543 469544 469543 469544 469546 469546 469546 469546 469546
BAC Name F18L15 F18L15 F18L15 F18L15	F18L13 F12A12 F12A12 F12A12 F12A12 F12A12	F12A12 F12A12 F12A12 F12A12 F12A12 F12A12 F12A12 F12A12 F12A12 F12A12 F12A12 F12A12 F12A12 F12A12 F12A12 F12A12 F12A12 F12A12 F12A12 F12A12
BAC Chromosome Length 3 100328 3 100328 3 100328	100328 100815 100815 100815 100815 100815	100815 100815
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Seq id AL133298 AL133298 AL133298	AL133314 AL133314 AL133314 AL133314 AL133314 AL133314 AL133314	AL133314
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SNP Base Columbia/	Landsberg																	G/T	G/C	C/T	A/T	G/A	C/G	A/G	A/G	A/C	A/G	T/A	A/T	C/A	G/A	T/C	T/G	CT	CT	C/A	T/C
Indel Size Columbia/	Landsberg	12/-12	-3/3	-3/3	-12/12	12/-12	9/9-	6-/6	-3/3	9-/9	4/-4	-1/1	-1/1	-2/2	-1/1	-1/1	-1/1																				
	Method	2	2	2	2	2	7	2	2	2	2			1	_	-	1			П	_	-	-		T	П					1	_	_	_		<b>T</b>	<b></b>
	Type	ONI	IND	IND	ON N	ONI	IND	ONI	ONI	ONI	ONI	ONI	IND		ONI	ONI	ONI	SNP	SNP																		
	Right	22621	22882	65752	73380	74891	75009	76933	83192	98928	99046	25824	25825	38422	61742	62185	65582	63615	64943	65437	64859	50872	50805	115576	38213	29722	29666	48696	48729	42915	42905	42318	42373	42321	42928	104101	104173
	Left	22608	22881	65751	73379	74878	75008	76923	83191	98921	99041	25823	25824	38421	61741	62184	65581	63613	64941	65435	64857	50870	50803	115574	38211	29720	29664	48694	48727	42913	42903	42316	42371	42319	42926	104099	104171
Marker	Name	469552	469553	469554	469555	469556	469557	469558	469559	469560	469561	471090	471091	471092	471093	471094	471095	467076	467319	467320	467321	467385	467386	467387	467985	468877	468878	469027	469028	469029	469030	469031	469032	469033	469034	469256	469257
BAC	Name	F12A12	F13112	F13I12	F13112	F13112	F13112	F13112	F13112		F13I12	F13112	$\overline{}$	F13112																							
BAC	Chromosome Length	100815	100815	100815	100815	100815	100815	100815	100815	100815	100815	100815	100815	100815	100815	100815	100815	140680	140680	140680	140680	140680	140680	140680	140680	140680	140680	140680	140680	140680	140680	140680	140680	140680	140680	140680	140680
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	Seq id	AL133314	AL133292	AL133292																																	
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SNP Base Columbia/ Landsberg A/G T/C	A/G G/A A/G A/G G/A G/A C/T
Indel Size Columbia/ Landsberg 12/-12 3/-3 -8/8 18/-18 -9/9 460/-460 12/-12 -14/14 2127/-2127 -12/12 3/-3 6/-6 3/-3 26/-26 14/-14 -8/8 -3/3 14/-14 -1/1 1/-1	
Method 1	
	SNP SNP SNP SNP SNP SNP SNP SNP SNP SNP
Right 104170 27492 107053 107097 109105 11478 11496 128221 13752 13752 13752 13754 18706 29668 47343 57439 57834 57834 57834 57834 57838 5	75144 7388 7162 7325 7485 62678 24860 59829 59534 59731 59745
Left 104168 27490 107040 107093 109104 11459 11495 127760 13739 13736 16578 29667 47339 57432 57830 88714 77616 80498 81265 82044 115562 27483 50364	55120 75142 7386 7160 7323 7483 62676 24858 59827 59729 59729 59743
Marker Name 469258 469467 469592 469593 469593 469595 469596 469599 469600 469600 469600 469600 469600 469600 469600 469600 469600 469600 469600 469600 469600 469600	4/1110 466981 467006 467008 467291 467291 467291 468339 468340 468341 468342
BAC Name F13112	F13112 F1P2 F1P2 F1P2 F1P2 F1P2 F1P2 F1P2 F
BAC Chromosome Length 140680 3 140680 3 140680	140880 101154 101154 101154 101154 101154 101154 101154 101154 101154
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Seq id AL133292 AL133292 AL133292 AL133292 AL133292 AL133292 AL133292 AL133292 AL133292 AL133292 AL133292 AL133292 AL133292 AL133292 AL133292 AL133292 AL133292 AL133292	AL133292 AL132955 AL132955 AL132955 AL132955 AL132955 AL132955 AL132955 AL132955 AL132955 AL132955 AL132955
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0, 0 H 41, 0 1, 1, 0 0 1	AVC C/G
Indel Size Columbia/ Landsberg 10/-10 -39/39 45/-45 4/-4 7/-7 3/-3 -3/3 -3/3 -3/3 -4/4 -10/10 3928/-3928 -15/15 -4/4 -1/-1 1/-1 1/-1 1/-1 -1/1 -1/1 -1/	
Method 1 1 1 1 1 2 2 2 2 2 2 2 2 2 2 2 2 2 2 2	<b>-</b>
	SNP
Right 914 82102 81922 82515 81922 82515 81969 81931 82117 25982 26188 51668 35571 70894 71915 72425 73702 81202 81483 82153 84082 84755 93649 94964 25965 59573 59573 59573	40336 36949
Left 912 82100 81920 82513 81920 82513 81929 82115 25980 26186 51666 35560 36346 66111 73701 81482 82152 84754 87825 89198 93648 94963 25963 59569 59571 59708	40334 36947
Marker Name 468449 468744 468745 468746 468747 468749 468749 469874 469845 469845 469845 469845 469845 469855 469855 469855 469857 469858	467027 467118
BAC Name FIP2 FIP2 FIP2 FIP2 FIP2 FIP2 FIP2 FIP2	T8P19 T8P19
BAC Chromosome Length 3 101154 3 101154	96679 96679
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Seq id AL132955	AL133315 AL133315
Seq # # # # # # # # # # # # # # # # # # #	85 85

SNP Base Columbia/	Landsberg	A/T	G/A	G/A	A/T	A/T	G/C	G/C	G/A	A/T	G/A	C/T	G/T	T/G	C/A	G/A	T/C	A/G	C/G	CT	G/T	A/T	A/G														
Indel Size Columbia/	Landsberg																							3/-3	L/L-	-3/3	-14/14	1145/-1145	39/-39	24/-24	8/8-	1639/-1639	1642/-1642	-15/15	-15/15	4/-4	772/-772
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	Left	36190	29081	29064	46757	47732	1105	1199	26769	26874	5481	76614	64659	67751	50477	50223	50392	50428	50355	50489	50424	50420	48413	12516	30508	36552	496	54336	6346	64844	72586	72778	73035	83589	83590	88118	90816
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SNP Base Columbia/	Landsberg					A/G	G/C	T/G	A/G	T/G	C/T	G/T	C/T	C/A	C/A	G/A	T/C	A/C	CT	A/T	C/T	T/A	T/A	G/A	G/A	T/A	T/C	T/C	T/C	G/C	G/C	T/G	A/G	1/G	A/G	D/O	A/G
Indel Size Columbia/	Landsberg	-1/1	-1/1	-1/1	-1/1																																
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	Type	ONI	IND		ON	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP
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	Left	1162	12259	29078	29081	25733	1513	1583	1576	1330	1581	1560	1143	24057	24045	40917	41278	41405	41088	41493	40214	42713	43277	41635	41644	41646	42323	43324	41623	41634	41645	42242	42688	43135	41636	41637	41639
Marker	Name	471622	471623	471624	471625	467175	467235	467236	467237	467238	467239	467240	467241	467406	467407	467451	467452	467453	467454	467455	467456	467457	467458	467459	467460	467461	467462	467463	467464	467465	467466	467467	467468	467469	467470	467471	467472
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	S	Э	$\varepsilon$	ĸ	$\epsilon$	$\mathfrak{C}$	3	E	$\varepsilon$	3	$\varepsilon$	$\epsilon$	ĸ	n	$\mathcal{E}$	3	$\epsilon$	$\epsilon$	$\mathfrak{C}$	3	3	$\varepsilon$	$\varepsilon$	3	$\mathfrak{C}$	3	$\varepsilon$	3	n	$\epsilon$	$\epsilon$	33	3	$\epsilon$	$\kappa$	$\kappa$	3
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SNP Base Columbia/	Landsberg	A/G	C/T	C/T	A/T	A/G	C/A	T/C	A/C	A/G	T/G	CT	G/A	T/A	T/C	C/G	G/C	T/C	$^{\mathrm{C/I}}$	A/G	C/T	C/T	T/A	T/A	G/A	C/G											
Indel Size Columbia/	Landsberg																										4/-4	-3/3	-15/15	L-1L	-74/74	8-/8	293/-293	772/-772	3223/-3223	11/-11	-3/3
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	Type	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	ONI		QNI	ONI	IND	ONI	ONI	ONI	ONI	ONI	ONI
	Right	41652	43280	41632	41634	19861	80963	77067	78887	79804	79217	79890	6761	78421	78174	78052	51261	53457	4619	92669	21454	22632	45149	48114	47977	46759	1007	25100	26739	27287	27639	33025	33928	4473	41674	38971	44921
	Left	41650	43278	41630	41632	19859	80961	79075	78885	79802	79215	79888	6229	78419	78172	78050	51259	53455	4617	67974	21452	22630	45147	48112	47975	46757	1002	25099	26738	27279	27638	33016	33634	3700	38450	38959	44920
Marker	Name	467473	467474	467475	467476	467594	467607	467608	467609	467610	467611	467612	467679	468254	468255	468256	468795	469005	469025	469224	469340	469436	469452	469453	469454	469455	470746	470747	470748	470749	470750	470751	470752	470753	470754	470755	470756
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BAC	Chromosome Length	82912	82912	82912	82912	82912	82912	82912	82912	82912	82912	82912	82912	82912	82912	82912	82912	82912	82912	82912	82912	82912	82912	82912	82912	82912	82912	82912	82912	82912	82912	82912	82912	82912	82912	82912	82912
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SNP Base Columbia/ Landsberg  Landsberg  T/A A/C G/A G/A A/G A/T A/G A/T	5
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Method	- 0 0 0 0
Right 49790 56190 57022 57909 59431 63824 65697 65493 65505 75928 1554 19925 21672 24060 24112 41310 41629 51369 6395 6678 81740 27929 28645 7701 7701 7701 8139 56683	22510 1118 18108 18110 25667
Left 49743 56289 57903 57903 57903 57903 63819 65096 65492 65504 75921 19827 19827 19827 19837 1987 1987 1987 1987 1987 1987 1987 198	32510 1034 18107 18109 25666
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SNP Base Columbia/	Landsberg					G/A	A/G	T/C	C/A	A/T	A/G	T/A	C/A	G/A	T/C	T/C	G/C	T/C	A/G	A/G	T/G	C/T	$C/\Gamma$	G/A	C/A	A/T	C/I	I/C	J/G	$C/\Gamma$	A/C	A/G	A/G	A/T	G/T	C/I	T/A
Indel Size Columbia/	Landsberg	3/-3	-17/17	-10/10	-1/1																																
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	Left	26019	46627	64104	27890	80963	80881	30883	19860	32445	26057	125484	126398	126201	125964	125616	126347	126244	125607	125574	125511	125573	125551	52591	52633	52642	52729	53602	53588	53763	54385	55381	54782	54541	54652	54829	57184
Marker	Name	470921	470922	470923	471565	466930	466931	466933	467117	467366	467550	467551	467552	467553	467554	467555	467556	467557	467558	467559	467560	467561	467562	467692	467693	467694	467695	467917	467918	467919	468194	468195	468196	468197	468198	468199	468200
BAC	Name	T2J13	T2J13	T2J13	T2J13	F2K15	F2K15	F2K15	F2K15	F2K15	F2K15	F2K15	F2K15	F2K15	F2K15	F2K15	F2K15	F2K15	F2K15	F2K15	F2K15	F2K15	F2K15	F2K15	F2K15	F2K15	F2K15	F2K15	F2K15	F2K15	F2K15	F2K15	F2K15	F2K15	F2K15	F2K15	F2K15
BAC	Chromosome Length	85109	85109	85109	85109	129757	129757	129757	129757	129757	129757	129757	129757	129757	129757	129757	129757	129757	129757	129757	129757	129757	129757	129757	129757	129757	129757	129757	129757	129757	129757	129757	129757	129757	129757	129757	129757
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SNP Base Columbia/	Landsberg	C/A	G/A	G/A	G/A	T/C	T/C	A/G	A/G	C/G	C/G	CT	G/T	C/T	T/C	T/C																					
Indel Size Columbia/	Landsberg																8-/8	8/8-	-10/10	-10/10	9-/9	8/8-	4/-4	83/-83	6-/6	-3/3	-3/3	L/L-	L/L-	L/L-	24/-24	87/-87	17/-17	9/9-	-13/13	3/-3	-1/1
	Method	-		1		1	_	_	_		1	_			T		2	2	2	2	2	2	2	2	2	2	2	2	2	2	2	2	2	2	2	2	<del></del>
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	Left	26800	26767	57851	57850	56449	57526	57355	57349	57727	57489	58036	57533	57378	63622	58812	102723	103802	111866	111871	114547	123467	126580	20970	27464	27508	27510	28037	3978	3980	61182	70283	72111	85070	85753	99225	52126
Marker	Name	468201	468202	468203	468204	468205	468206	468207	468208	468209	468210	468211	468212	468213	468315	468520	470165	470166	470167	470168	470169	470170	470171	470172	470173	470174	470175	470176	470177	470178	470179	470180	470181	470182	470183	470184	471301
BAC	Name	F2K15																																			
BAC	Chromosome Length	3 129757	3 129757	3 129757	3 129757	3 129757	3 129757	3 129757	3 129757	3 129757	3 129757	3 129757	3 129757	3 129757	3 129757	3 129757	3 129757	3 129757	3 129757	3 129757	3 129757	3 129757				3 129757			3 129757	3 129757	3 129757	1			3 129757	3 129757	3 129757
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Indel Size Columbia/	Landsberg	-1/1																																			
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	Right	80720	51632	50492	50383	52327	52334	51626	52403	52366	52413	58480	57623	56669	69091	44208	43976	45746	45700	45865	55442	55958	53907	93559	93385	93624	22897	100661	100431	100799	100731	100980	14578	15442	81748	38079	36818
	Left	80719	51630	50490	50381	52325	52332	51624	52401	52364	52411	58478	57621	26667	68069	44206	43974	45744	45698	45863	55440	55956	53905	93557	93383	93622	22895	100659	100429	100797	100729	100978	14576	15440	81746	38077	36816
Marker	Name	471302	467146	467147	467148	467149	467150	467151	467152	467153	467154	467525	467526	467527	467701	468145	468146	468521	468522	468523	468787	468788	468789	468859	468860	468861	468868	468981	468982	468983	468984	468985	469016	469017	469290	469310	469345
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BAC	Chromosome Length	129757	104204	104204	104204	104204	104204	104204	104204	104204	104204	104204	104204	104204	104204	104204	104204	104204	104204	104204	104204	104204	104204	104204	104204	104204	104204	104204	104204	104204	104204	104204	104204	104204	104204	104204	104204
	Chr	$\mathcal{C}$	33	$\mathcal{C}$	3	3	3	3	3	3	3	$\epsilon$	3	3	3	3	3	$\epsilon$	$\mathfrak{C}$	3	$\mathcal{C}$	$\epsilon$	n	$\varepsilon$	$\epsilon$	$\mathfrak{C}$	$\epsilon$	$\epsilon$	$\epsilon$	$\epsilon$	$\epsilon$	$\mathcal{C}$	m	3	$\mathcal{C}$	8	$\alpha$
	Seq id	AL132956	AL132964	AL132964	AL132964	AL132964	AL132964	AL132964	AL132964	AL132964	AL132964	AL132964	AL132964	AL132964	AL132964	AL132964	AL132964	AL132964	AL132964	AL132964	AL132964	AL132964	AL132964	AL132964	AL132964	AL132964	AL132964	AL132964	AL132964	AL132964	AL132964	AL132964	AL132964	AL132964	AL132964	AL132964	AL132964
Sed	umu	88	68	68	68	68	68	68	68	68	68	68	68	68	68	68	68	68	68	68	68	68	68	68	68	68	68	68	68	68	68	68	68	68	68	68	68

SNP Base Columbia/ Landsberg G/A T/C	<b>4</b> 5
Indel Size Columbia/ Landsberg -7/7 6/-6 -222/22 -3/3 -5/5 -3/3 -4/4 -5/5 -17/17 -6/6 4/-4 -3/3 8/-8 -3/3 8/-8 -3/3 8/-8 -1/1 -1/-1 1/-1 1/-1 1/-1 1/-1 1/-	
Method	7
	SNP
Right 47428 47043 12101 16218 1856 2486 33581 35681 45129 48191 48816 51372 62721 66305 68120 70841 7684 14311 15163 15242 47653 50391 51376 51376 51653 69057 80494 93375	64402
Left 47426 47041 12100 16211 1855 2485 33580 35680 45128 4815 51371 62716 66304 68119 70840 70840 7387 76027 76579 9930 101152 14308 15162 15241 47620 47651 50389 51375 51375 69055 80492 93373	64400
Marker Name 469428 469429 471056 471056 471059 471060 471060 471061 471062 471063 471073 471073 471073 471073 471629 471639 471633 471639 471633 471635 471636 471639	466811
BAC Name 1963 1963 1963 1963 1963 1963 1963 1963	T16K5
BAC Chromosome Length 104204 3 104204 3 104204	97711
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Seq id AL132964	AL132965
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SNP Base Columbia/	Landsberg	A/G	A/C	T/G	G/T	C/I	G/A	G/A	T/A	A/C	CT	T/C	A/C														T/C	C/I	C/T	T/C	T/G	G/A	G/A	G/A	T/G	CA	C/I
Indel Size Columbia/	Landsberg													4/-4	-3/3	2/-2	-10/10	-4/4	6-/6	-3/3	1/-1	1/-1	3/-3	2/-2	1/-1	1/-1											
	Method	1	1	П	_	_			_	1	_	_	_	2	2	2	2	2	2	2	-	-		1	П	П	<b>—</b>			1	1	-		1			T
	Type	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	ONI	ONI	ONI		ONI	ONI	ONI	ONI	ONI	ONI	ONI	ON N	ONI	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP
	Right	64640	63375	29997	76920	75840	55187	41746	89190	89023	89301	77382	12896	50463	57949	64274	70596	79148	96240	96288	17463	30292	64275	64278	76523	76537	70277	11607	11684	10353	11022	67516	65400	59633	59473	41319	41243
	Left	64638	63373	99992	76918	75838	55185	41744	89188	89021	89299	77380	12894	50458	57948	64268	70595	79147	96230	96287	17461	30290	64271	64275	76521	76535	70275	11605	11682	10351	11020	67514	65398	59631	59471	41317	41241
Marker	Name	466812	466813	467542	467543	467544	467816	468158	468524	468525	468526	468929	468959	470613	470614	470615	470616	470617	470618	470619	471439	471440	471441	471442	471443	471444	467653	467673	467674	467675	467676	467722	468265	468309	468310	468364	468365
BAC	Name	T16K5	T16K5	T16K5	T16K5	T16K5	T16K5	T16K5	T16K5	T16K5	T16K5	T16K5	T16K5	T16K5	T16K5	T16K5	T16K5	T16K5	T16K5	T16K5	T16K5	T16K5	T16K5	T16K5	T16K5	T16K5	F3A4	F3A4	F3A4	F3A4	F3A4	F3A4	F3A4	F3A4	F3A4	F3A4	F3A4
BAC	Chromosome Length	97711	97711	97711	97711	97711	97711	97711	97711	97711	97711	97711	97711	97711	97711	97711	97711	97711	97711	97711	97711	97711	97711	97711	97711	97711	108158	108158	108158	108158	108158	108158	108158	108158	108158	108158	108158
	Chrc	$\mathcal{C}$	$\varepsilon$	$\mathfrak{C}$	3	3	$\kappa$	33	3	æ	33	3	$\epsilon$	3	33	æ	3	$\epsilon$	3	$\epsilon$	3	3	33	3	3	3	3	3	3	3	$\varepsilon$	$\mathfrak{C}$	$\varepsilon$	3	3	33	33
	Seq id	AL132965	AL132965	AL132965	AL132965	AL132965	AL132965	AL132965	AL132965	AL132965	AL132965	AL132965	AL132965	AL132965	AL132965	AL132965	AL132965	AL132965	AL132965	AL132965	AL132965	AL132965	AL132965	AL132965	AL132965	AL132965	AL132978	AL132978	AL132978	AL132978	AL132978	AL132978	AL132978	AL132978	AL132978	AL132978	AL132978
Seq	unu	96	90	90	06	90	90	90	90	90	90	8	90	90	8	8	06	90	06	90	06	8	90	06	06	06	16	91	91	91	91	91	91	91	91	91	91

SNP Base Columbia/ Landsberg G/A T//A C/T	A/G A/T 1/A 1/C G/T A/T
Indel Size Columbia/ Landsberg 3/-3 -8/8 -8/8 8/-8 10/-10 12/-12 3/-3 -11/11 -3/3 -1/1 -1/1 -1/1 -1/1 -1	
f Method	
	SNP SNP SNP SNP SNP SNP
Right 7803 64194 52392 14682 14768 14769 15995 18026 18113 26576 26624 3808 3856 56634 66813 8394 90535 92209 10793 29730 29807 7506 7515 8395	65945 51719 95293 35979 75596 74538 76703
Left 7801 64192 52390 14678 14767 14768 15986 18015 18100 266572 26623 3798 3855 56662 66805 8393 90509 92208 10792 29729 29729 7505 7514 8394	65943 51717 95291 35977 75594 74536 76701
Marker Name 468388 468642 469364 470200 470200 470202 470203 470203 470203 470203 470204 470209 470210 470211 470212 470213 470213 471309 471313	466904 467535 467615 467644 467822 467823 467824
BAC Name F3A4 F3A4 F3A4 F3A4 F3A4 F3A4 F3A4 F3A4	F11C1 F11C1 F11C1 F11C1 F11C1 F11C1
BAC Chromosome Length  108158	105644 105644 105644 105644 105644 105644
H O m m m m m m m m m m m m m m m m m m m	
Seq id AL132978	AL132976 AL132976 AL132976 AL132976 AL132976 AL132976 AL132976 AL132976
Seq m m 2 1 1 2 2 2 2 2 2 2 2 2 2 2 2 2 2 2 2 2	5 5 5 5 5 5 5 5 5 5 5 5 5 5 5 5 5 5 5

SNP Base Columbia/	Landsberg	G/A	T/A	A/C	A/G	A/G	T/C	A/G	A/T	G/A	A/G	T/A	T/C	T/C	CT	G/T	CT	A/T	CT	A/T	D/L	C/T	T/C	T/A	G/T	T/C											
Indel Size Columbia/	Landsberg	1																									9/9-	4/-4	-24/24	8/8-	10477/-	104//	1//-1/	1.1-1.1	9/9-	4/-4	-11/11
	Method	П	1	-			1	<b>,</b>						<del></del>			_				_	1	1	1			2	2	2	2	2	ď	7 (	7	7	7	2
	Type	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	IND	IND	IND	ONI	ONI	í	ONI Civi	ON!		ON!													
	Right	71969	73219	73386	72515	34352	91405	91420	91210	82992	80122	103050	103377	102734	102705	102717	102737	102795	103095	43124	38830	45856	40592	94341	94538	99014	101584	101921	104312	105094	12962	0	30337	30/8/	30773	47465	47578
	Left	71967	73217	73384	72513	34350	91403	91418	91208	82990	80120	103048	103375	102732	102703	102715	102735	102793	103093	43122	38828	45854	40590	94339	94536	99012	101583	101916	104311	105093	2484	0	30519	30/69	30772	47460	47577
Marker	Name	467826	467827	467828	467829	467934	467971	467972	467973	468078	468118	468536	468537	468538	468539	468540	468541	468542	468543	468796	469231	469232	469233	469235	469236	469411	469494	469495	469496	469497	469498	00,000	469499	469500	469501	469502	469503
BAC	Name	F11C1	FIICI	FIICI	F11C1	FIICI	F11C1	F11C1	F11C1	F11C1	F11C1	F11C1	F11C1	F11C1	F11C1	F11C1	F11C1	F11C1	F11C1	F11C1	F11C1	FIICI	F11C1	F11C1	F11C1	Ç	FIICI	FIICI	F11C1	FIICI	F11C1						
BAC	Chromosome Length	105644	105644	105644	105644	105644	105644	105644	105644	105644	105644	105644	105644	105644	105644	105644	105644	105644	105644	105644	105644	105644	105644	105644	105644	105644	105644	105644	105644	105644	105644	100	105644	105644	105644	105644	105644
	Chromo	3	3	3	3	3	3	3	3	3	3	3	3	3	3	3	3	3	3	3	3	3	3	3	3	3	3	3	3	3	3	,	<b>n</b> (		m .	ε,	m
	Seq id	AL132976	AL132976	AL132976	AL132976	AL132976	AL132976	AL132976	AL132976	AL132976	AL132976	AL132976	AL132976	AL132976	AL132976	AL132976	AL132976	AL132976	AL132976	70000114	AL132970	AL1329/6	AL132976	AL132976	AL132976												
Seq	unu	92	92	35	92	92	92	95	92	92	35	95	92	95	92	95	92	92	95	65	65	92	92	92	92	92	92	92	92	92	92	5	76	76	92	92	92

SNP Base Columbia/ Landsberg  T/C C/T A/G C/T T/A T/A T/A G/A A/G C/T T/A T/A T/G A/G C/T
Indel Size Columbia/ Landsberg 6/-6 4/-4 6/-6 -14/14 -5/5 -1/1 1/-1 1/-1 1/-1 1/-1 1/-1 1/-1
0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0
Right 53718 53718 64844 64869 69148 97569 14671 34297 67142 71272 73371 73371 73373 73907 94139 12173 55248 67770 6702 5524 6401 5524 6401 5524 6401 5524 6401 12173 53248 67770 73520 6702 5702 45934 41575 42701 11229 11326 11326
Left 53711 53711 53713 64862 69147 97568 14670 34296 67140 71271 73369 73371 73906 94138 12171 52932 53015 53246 6700 5522 6399 5843 7200 45932 41573 42699 42741 42699 42741 11222 11322
Marker Name 469504 469504 469505 469506 469507 469508 469509 471082 471083 471084 471088 467035 471088 467035 471088 467035 471089 467035 467035 467035 467035 467035 467035 467035 467035 468799 468799 468799 468799 468799 468799 468799 468799 468799 468798 468799 468799 468799 468799 468799 468799 468799 468799 468799 468799 468799
BAC Name F11C1 F11C2 F11
BAC 105644 106644 10664 106644 10664
Chromosome
Seq id AL132976 AL133363
8 2 3 2 3 2 3 2 3 2 3 2 3 2 3 2 3 2 3 2

SNP Base Columbia/ Landsberg	G/A 17/C 17/C 17/C 17/C 17/C 17/C 17/C 17/C
Indel Size Columbia/ Landsberg 3/-3 -3/3 -24/24 -22/22 2/-2 -1/1 -1/1 4/-4 -1/1	4/-4 16/-16 10/-10 1016/-1016 3/-3 5/-5 3/-3 6/-6 -6/6 1004/-1004
Method 2 2 2 1 1 1 1	
Type CNI	SY S
Right 27935 28583 64837 8688 42082 42988 43301 45152 53066 53249 5622	20016 19735 26957 46424 57882 16667 16803 62498 18190 17894 70358 41142 14916 11658 11944 13540 43756 45972 51998 5848 65524
Left 27931 28582 64836 8687 42079 42987 43300 45147 53065 55224	20014 20014 19733 26955 46422 57880 16665 16801 62496 18188 17892 70356 41140 14914 11953 11927 13529 42739 42739 65523 76181
Marker Name 470722 470723 470724 471484 471485 471485 471486 471489 471489	47,1490 467,144 467,145 467,311 468076 468077 468181 468358 468358 468358 468357 468357 468357 469341 470944 470945 470945 470950 470953 470953
BAC Name T20E23 T20E23 T20E23 T20E23 T20E23 T20E23 T20E23 T20E23 T20E23	T20E23 T3A5 T3A5 T3A5 T3A5 T3A5 T3A5 T3A5 T3A
BAC Chromosome Length 3 83513 3 83513 3 83513 3 83513 3 83513 3 83513 3 83513 3 83513 3 83513 3 83513	83513 84196 84196 84196 84196 84196 84196 84196 84196 84196 84196 84196 84196 84196 84196 84196 84196
Chrc 33 33 33 33 33 33 33 33 33 33 33 33 33	
Seq id AL133363 AL133363 AL133363 AL133363 AL133363 AL133363 AL133363 AL133363 AL133363	AL133363 AL132979 AL132979 AL132979 AL132979 AL132979 AL132979 AL132979 AL132979 AL132979 AL132979 AL132979 AL132979 AL132979 AL132979 AL132979 AL132979 AL132979 AL132979 AL132979
Seq 93 93 93 93 93 93	2

SNP Base Columbia/ Landsberg	1/G A/G A/T G/T G/A	1/C 1/C 1/C C/T	A/T G/A 1/G G/T C/G	AC C/T C/T C/T A/G 1/A A/G
Indel Size Columbia/ Landsberg -6/6 -1/1 1/-1 3/-2 -1/1	}			
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Type GNI GNI GNI GNI GNI GNI GNI GNI GNI GNI	SNP SNP SNP SNP SNP SNP	SNP SNP SNP SNP SNP	SNP SNP SNP SNP SNP SNP SNP SNP SNP SNP	S S S S S S S S S S S S S S S S S S S
Right 78015 14994 40956 45985 46621	102171 101970 102144 78439 84797 84812	84806 84795 84791 84815 84509	84441 37400 18926 18811 123953 123730 100642 100632 26309	11403 11833 13036 12995 12784 13475 62113 67145 67362 66900
Left 78014 14993 40954 45981 46618 62639	102169 101968 102142 78437 84795 84810	84793 84793 84789 84813 84507	84439 37398 18924 18809 123951 123728 100640 100630 26307	11401 11831 13034 12993 12782 13473 62111 67143 66898
Marker Name 470955 471574 471575 471576 471577	466826 466827 466828 467166 467186	467188 467189 467190 467191	467193 467256 467257 467258 467907 467908 468017 468018	468270 468271 468273 468273 468275 468375 46869 468670
BAC Name T3A5 T3A5 T3A5 T3A5 T3A5	F24M12 F24M12 F24M12 F24M12 F24M12 F24M12	F24M12 F24M12 F24M12 F24M12 F24M12	F24M12 F24M12 F24M12 F24M12 F24M12 F24M12 F24M12	F24M12 F24M12 F24M12 F24M12 F24M12 F24M12 F24M12 F24M12
BAC Chromosome Length 3 84196 3 84196 3 84196 3 84196 3 84196	129516 129516 129516 129516 129516 129516	129516 129516 129516 129516 129516	129516 129516 129516 129516 129516 129516 129516 129516	129516 129516 129516 129516 129516 129516 129516 129516
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Seq id AL132979 AL132979 AL132979 AL132979	AL132980 AL132980 AL132980 AL132980 AL132980 AL132980	AL132980 AL132980 AL132980 AL132980 AL132980	AL132980 AL132980 AL132980 AL132980 AL132980 AL132980 AL132980	AL132980 AL132980 AL132980 AL132980 AL132980 AL132980 AL132980 AL132980 AL132980
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SNP Base Columbia/ Landsberg	A/G	T/G	C/T	CT	C/A	G/A	C/T	T/C	A/G	G/A	G/A	T/A	T/G	A/G	C/A	A/C	G/A																			
Indel Size Columbia/ Landsberg	0																	11/-11	-24/24	14/-14	14/-14	-4/4	10/-10	-62/62	13/-13	29/-29	-10/10	94/-94	8-/8	-4/4	-72/72	-3/3	-5/5	-2/2	1/-1	1/-1
Method	1	-	-	1					_			_	, <b>-1</b>	<b>—</b>	_	_	_	2	2	2	7	2	2	2	2	2	2	2	2	2	2	2	2	1	-	
Tvne	SNP	SNP	SNP	SNP	SNP	ONI	ON.	ONI	ONI		ONI	ONI	ONI		QNI				ONI			ONI	ON ON	QN N												
Right	67138	67082	67563	22029	32319	32269	32140	94561	94142	5859	5862	5933	6421	5932	9644	9587	97881	108809	111360	116751	116776	127478	128402	14335	40140	43518	52984	70651	72467	75637	0/09/	82862	98401	18728	37232	9182
Left	67136	67080	67561	67075	32317	32267	32138	94559	94140	5857	2860	5931	6419	5930	9642	9585	61816	108797	111359	116736	116761	127477	128391	14334	40126	43488	52983	70556	72458	75636	69092	82861	98400	18727	37230	9180
Marker Name	468672	468673	468674	468675	468898	468899	468900	468937	468938	469102	469103	469104	469105	469106	469114	469115	469391	469953	469954	469955	469956	469957	469958	469959	469960	469961	469962	469963	469964	469965	469966	469967	469968	471235	471236	471237
BAC Name	F24M12	F24M12	F24M12	F24M12	F24M12	F24M12	F24M12	F24M12	F24M12	F24M12	F24M12	F24M12	F24M12	F24M12	F24M12	F24M12	F24M12	F24M12	F24M12	F24M12	F24M12	F24M12	F24M12	F24M12												
BAC some Length		129516	129516	129516	129516	129516	129516	129516	129516	129516	129516	129516	129516	129516	129516	129516	129516	129516	129516	129516	129516	129516	129516	129516	129516	129516	129516	129516	129516	129516	129516	129516	129516	129516	129516	129516
Chromosome	3	3	3	3	3	3	3	3	3	3	Э	3	Э	33	3	3	3	Э	3	3	3	3	3	3	3	3	33	3	3	$\mathcal{E}$	3	3	3	3	$\mathcal{C}$	33
Seo id	AL132980	AL132980	AL132980	AL132980	AL132980	AL132980	AL132980	AL132980	AL132980	AL132980	AL132980	AL132980	AL132980	AL132980	AL132980	AL132980	AL132980	AL132980	AL132980	AL132980	AL132980	AL132980	AL132980	AL132980												
Seq	95	95	95	95	95	95	95	95	95	95	95	95	95	95	95	95	95	95	95	95	95	95	95	95	95	95	95	95	95	95	95	95	95	95	95	95

SNP Base Columbia/ Landsberg	T/C	T/C	T/C	G/A																		C/A	T/A	T/A	A/G	G/T	G/C	T/A	C/A	G/C	T/C	A/G	A/G	A/G	G/A
Indel Size Columbia/ Landsberg	-2/2				-4/4	10/-10	9-/9	5036/-5036	14/-14	14/-14	-12/12	8/8-	6-/6	6-/6	8/8-	9-/9	8/8-	18/-18	18/-18	4/-4	3/-3														
Method		1	1	1	7	2	2	2	2	2	2	2	2	2	2	2	2	2	2	2	2	1	_	-	-	_			1	_	1		1	1	
Type	ONI SNP	SNP	SNP	SNP	ONI	IND	IND	ON.	ONI	IND	ONI	ONI	IND	ON.	ONI	ONI	ONI	ONI	ONI	IND	ONI	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP
Right	9206 64271	54009	12617	15989	13874	14798	28425	35715	3147	3172	32199	33869	52395	52402	56629	66239	66673	77465	77486	83534	86906	65188	65377	65378	66244	66176	17741	49061	49681	49059	49233	49305	61328	61342	21986
Left	9205 64269	54007	12615	15987	13873	14787	28418	30678	3132	3157	32198	33868	52385	52392	56628	66232	66672	77446	77467	83529	90694	65186	65375	65376	66242	66174	17739	49059	49679	49057	49231	49303	61326	61340	57984
Marker Name	471238 467578	467920	468350	468620	470030	470031	470032	470033	470034	470035	470036	470037	470038	470039	470040	470041	470042	470043	470044	470045	470046	467242	467243	467244	467245	467246	467598	467664	467665	467666	467667	467668	467761	467762	467763
BAC Name	F24M12 F26O13	F26013	F26013	F26013	F26013	F26013	F26013	F26013	F26013	F26013	F26013	F26013	F26013	F26013	F26013	F26013	F26013	F26013	F26013	F26013	F26013	T18N14	T18N14	T18N14	T18N14	T18N14	T18N14	T18N14	T18N14	T18N14	T18N14	T18N14	T18N14	T18N14	T18N14
BAC Chromosome Length	129 <u>5</u> 16 94349	94349	94349	94349	94349	94349	94349	94349	94349	94349	94349	94349	94349	94349	94349	94349	94349	94349	94349	94349	94349	L986L	19861	19861	19867	19861	19861	L986L	19861	19861	19861	19861	L986L	19861	19861
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Seq id	AL133980 AL133452	AL133452	AL133452	AL133452	AL133452	AL133452	AL133452	AL133452	AL133452	AL133452	AL133452	AL133452	AL133452	AL133452	AL133452	AL133452	AL133452	AL133452	AL133452	AL133452	AL133452	AL132968	AL132968	AL132968	AL132968	AL132968	AL132968	AL132968	AL132968	AL132968	AL132968	AL132968	AL132968	AL132968	AL132968
Seq	95	96	96	96	96	96	96	96	96	96	96	96	96	96	96	96	96	96	96	96	96	24	26	26	26	26	24	26	26	26	26	24	26	24	26

SNP Base Columbia/ Landsberg G/A T/C C/G G/T G/A A/T C/A T/A G/A G/T C/A T/A C/A T/A C/A T/A C/T C/A T/A C/T C/A T/A C/T C/A T/A C/T C/A C/T C/A C/T C/A C/T C/A C/T C/A C/T C/A C/T C/T C/A C/T	
Indel Size Columbia/ Landsberg Landsberg 18/-18 18/-18 6/-6 4/-4 -8/8 3/-3 -10/10	4/-4 -3/3 20/-20
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Right 57835 57941 57926 58205 19099 20011 19527 19263 20332 19456 63257 63439 30258 30258 30258 30258 30258 71134 71007 70222 71134 12965 1718 19013 2152 26177	48287 53499 60802
Left 57833 57939 57924 58203 19097 20009 19525 19261 20330 19454 63255 63437 30135 30257 29668 30087 24775 4810 28024 27942 56421 56421 5662 71132 12925 12946 1711 19008 2151	48282 53498 60781
Marker Name 467764 467765 467766 467767 468220 468221 468222 468223 468223 468224 468224 468224 468438 468439 468438 468438 468438 469438 469409 469080 469080 469335 469335 469335 469335 469469 470660 470663	470666 470667 470668
BAC Name T18N14	T18N14 T18N14 T18N14
Chromosome Length  3 3 19867	
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Seq num num 97 97 97 97 97 97 97 97 97 97 97 97 97	97

SNP Base Columbia/ Landsberg A/G T/A T/C C/G G/T A/C T/A	17.4 A/G G/T C/T 17.C C/G G/A 17.C
Indel Size Columbia/ Landsberg 5/-5 12/-12 12/-12 2/-2 -1/1 -1/1 -1/1 -1/1	ı i
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Right 65673 79319 79323 19011 19011 19014 24753 29620 29625 49151 49212 49225 63108 63173 65597 66173 32172 55868 56133 56113 65588 42265 18232 65397 65397 65397 65397 65397 65397 65455	13821 13821 13673 17230 67335 66593 209
Left 65667 79306 79310 19008 19011 24752 29624 49149 49210 49224 63172 63172 65594 66172 55866 56111 28406 18172 65343 65343 65343 65346 18231 65396	12607 13819 13671 17228 67333 66591 207
Marker Name 470669 470670 470671 471460 471462 471463 471464 471465 471466 471469 471469 471470 471470 471471 468165 468167	466797 466798 466799 466829 467141 467142 467159
BAC Name T18N14 T18N14 T18N14 T18N14 T18N14 T18N14 T18N14 T18N14 T18N14 T18N14 T18N14 T18N14 T18N14 T18N14 T18N15 T25B15 T25B15 T25B15 T25B15 T25B15 T25B15 T25B15 T25B15 T25B15 T25B15 T25B15 T25B15 T25B15 T25B15 T25B15	F812 F812 F812 F812 F812 F812 F812
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	Right	480	165	484	478	159	2619	2586	65530	7610	7585	47461	47778	47473	78474	90983	69620	42122	50419	50255	24365	56736	34517	19308	16130	16247	60868	89831	69268	73502	5487	5525	5837	5856	55294	94650	94570
	Left	478	163	482	476	157	2617	2584	65528	2097	7583	47459	47776	47471	78472	90981	69618	42120	50417	50253	24363	56734	34515	19306	16128	16245	89807	89829	19168	73500	5485	5523	5835	5854	55292	94648	94568
Marker	Name	467161	467162	467163	467164	467165	467196	467197	467230	467373	467374	467502	467503	467504	467563	467700	467958	467970	468131	468132	468313	468323	468329	468514	468604	468605	468727	468728	468729	469026	469054	469055	469056	469057	469116	469159	469160
BAC	Name	F8J2	F8J2	F8J2	F8J2	F8J2	F8J2	F8J2	F8J2	F8J2	F8J2	F8J2	F8J2	F8J2	F812	F8J2	F8J2	F8J2	F8J2	F8J2	F8J2	F8J2	F8J2	F8J2	F8J2	F8J2	F812	F8J2	F8J2	F8J2	F8J2	F812	F812	F812	F8J2	F812	F812
BAC	Chromosome Length	86176	3 97798	861798	861168	861168	3 97798	86176	861168	86176	867798	861168	867798	861798	3 97798	86176	3 97798	3 97798	3 97798	3 97798	3 97798	3 97798	3 97798	3 97798	86776	3 97798	3 97798	367798	3 97798	3 67798	3 97798	3 97798	3 97798	3 67798	3 97798	3 67798	3 97798
	Seq id (	AL132969 3		AL132969 3	AL132969 3	AL132969 3	AL132969 3	AL132969 3	AL132969 3	AL132969 3	AL132969 3	AL132969 3	AL132969	AL132969	AL132969 3	AL132969	AL132969 3	AL132969																			
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SNP Base Columbia/ Landsberg A/G G/T		ţ.	A/G A/G G/T A/T A/G C/G
Indel Size Columbia/ Landsberg 46/-46	40/-46 -9/9 -3/3 6/-6 6/-6 4/-4 9/-9 -6/6 -14/14 1/-1	1,-1 1,-1 1,-1 1,-1 1,-1 1,-1 1,-1 -1,1 -1,1 -2,2 2,-2	
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Right 94513 94608 94489 10694	10694 33794 35749 35830 52915 56623 77014 79525 81908	13310 18053 19473 23628 24183 24470 24694 50395 56626 67549 87729 89906 90862	47901 46955 74750 74824 74825 20880 20979 71120
Left 94511 94606 94487 10647	10647 33793 35748 35823 52908 56618 77004 79524 81907	13308 18051 18051 19472 23626 24181 24692 50394 56621 67548 87728 89905 90861	47899 46953 74748 74822 74823 20878 20977 71118
Marker Name 469161 469162 469163	470385 470386 470388 470389 470391 470392 470393 471358	471359 471360 471362 471363 471364 471365 471366 471369 471370 471373	466946 466947 467247 467248 467249 467576 467577
BAC Name F812 F812 F812 F812	F812 F812 F812 F812 F812 F812 F812 F812	F812 F812 F812 F812 F812 F812 F812 F812	14D2 14D2 14D2 14D2 14D2 14D2 14D2
BAC Chromosome Length 3 97798 3 97798 3 97798	97798 97798 97798 97798 97798 97798 97798	97798 97798 97798 97798 97798 97798 97798 97798	92611 92611 92611 92611 92611 92611
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	Right	70205	6229	6334	7959	45058	45053	45038	44627	45039	45026	45041	45019	44618	59568	77476	28384	28202	33655	34572	31908	34099	32546	64245	18496	61964	61883	55487	30622	30653	3295	40847	44202	51889	57131	69802	78971
	Left	70203	6227	6332	7957	45056	45051	45036	44625	45037	45024	45039	45017	44616	29566	77474	28382	28200	33653	34570	31906	34097	32544	64243	18494	61962	61881	55485	30621	30652	3264	40835	44201	51879	57125	70861	0968L
Marker	Name	467760	467817	467818	467819	467987	467988	467989	467990	467991	467992	467993	467994	467995	468229	468372	468678	468679	468719	468720	468721	468722	468723	468739	468814	469123	469124	469220	470970	470971	470972	470973	470974	470975	470976	470977	470978
BAC	Name	T4D2	T4D2	T4D2	T4D2	T4D2	T4D2	T4D2	T4D2	T4D2	T4D2	T4D2	T4D2	T4D2	T4D2	T4D2	T4D2	T4D2	T4D2	T4D2	T4D2	T4D2	T4D2	T4D2	T4D2	T4D2	T4D2	T4D2	T4D2	T4D2	T4D2	T4D2	T4D2	T4D2	T4D2	T4D2	T4D2
BAC	Chromosome Length	92611	92611	92611	92611	92611	92611	92611	92611	92611	92611	92611	92611	92611	92611	92611	92611	92611	92611	92611	92611	92611	92611	92611	92611	92611	92611	92611	92611	92611	92611	92611	92611	92611	92611	92611	92611
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	Seq id	AL132958	AL132958	AL132958	AL132958	AL132958	AL132958	AL132958	AL132958	AL132958	AL132958	AL132958	AL132958	AL132958	AL132958	AL132958	AL132958	AL132958	AL132958	AL132958	AL132958	AL132958	AL132958	AL132958	AL132958	AL132958	AL132958	AL132958	AL132958	AL132958	AL132958	AL132958	AL132958	AL132958	AL132958	AL132958	AL132958
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SNP Base Columbia/ Landsberg	G/A A/C A/G G/T T/G	A/G C/A A/G G/A	G/A G/A G/A G/A G/T A/T	A/T T/G G/C G/A C/A C/T
Indel Size Columbia/ Landsberg 10/-10 1/-1 1/-1 1/-1				3/-3 3/-3
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Type IND IND IND IND	SNP SNP SNP SNP	SNP SNP SNP SNP SNP	S S S S S S S S S S S S S S S S S S S	SNP
Right 78999 14671 20166 28156 55618	68996 73411 72362 46941 99696	104388 105634 59967 60316 41152 88922	66749 55469 55426 110036 110584 109642 80598 94927 65809	26595 10680 137904 137907 137903 28328 82945 51900 109657
Left 78988 14669 20164 28154 55615	68994 73409 72360 46939 99694	104386 105632 59965 60314 41150 88920	66747 55467 55424 110034 110582 109640 80596 94925 65807	26593 10678 137902 137905 137901 28326 82943 51940 51898 109653
Marker Name 470979 471581 471582 471583	466952 466953 466954 467098	467179 467180 467299 467300 467436	467497 467662 467663 467732 467733 467734 467874 467874	468253 468461 468480 468481 468482 468975 469175 469176 470278
BAC Name T4D2 T4D2 T4D2 T4D2 T4D2	F4P12 F4P12 F4P12 F4P12 F4P12	F4P12 F4P12 F4P12 F4P12 F4P12	F4P12 F4P12 F4P12 F4P12 F4P12 F4P12 F4P12 F4P12	F4P12 F4P12 F4P12 F4P12 F4P12 F4P12 F4P12 F4P12 F4P12
BAC Chromosome Length 3 92611 3 92611 3 92611 3 92611	144628 144628 144628 144628	144628 144628 144628 144628 144628	144628 144628 144628 144628 144628 144628 144628	144628 144628 144628 144628 144628 144628 144628 144628
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Columbia/ Landsberg T/C A/T	G/A
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Right 129956 138890 139203 20334 18795 18801 19833 19833 19835 29113 31998 32004 33319 38767 45710 46420 48152 49197 77219 77239 81423 109662 22361 28131 28140 65806 66709 72815 73027 73032 95069 106869 2246	68//6
Left 129946 133877 139202 17351 18774 18780 19804 19806 29112 31997 32003 33758 45709 46419 48146 49192 57398 77218 77218 77218 77218 77218 77218 77218 77218 77218 77218 77218 77218 77218 73019 77218 77229 66708 773029 95068	18//6
Marker Name 470280 470281 470282 470283 470283 470284 470284 470286 470293 470299 470299 470299 470299 470299 470299 470299 470399 470399 470399 470399 470399 471335 471335 471334 471334 471334 471334 471334	40707
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Indel Size Columbia/	Landsberg	•																										-15/15	L'IL-	3/-3	9/9-	-11/11	10/-10	-34/34	13/-13	-15/15	-15/15
	Method		1	1		_	1	1	1	1	_	П	=	ī	1	1		_	1	1	П	_		<b>-</b>	_	П		2	2	2	2	2	2	2	2	2	2
	Type	SNP	IND	IND	ONI	ONI	IND	ONI	ONI	ONI	ONI	ONI																									
	Right	57857	111607	111502	35460	30817	30934	30936	9598	85623	85124	85063	100416	8208	8057	8754	105430	10430	10542	25823	25089	20123	22903	90025	90213	43419	44006	101850	1042	107414	108513	109026	29740	47234	54173	61921	61926
	Left	57855	111605	111500	35458	30815	30932	30934	9656	85621	85122	85061	100414	8706	8055	8752	105428	10428	10540	25821	25087	20121	22901	90023	90211	43417	44004	101849	1041	107410	108512	109025	29729	47233	54159	61920	61925
Marker	Name	467510	467529	467530	467545	467566	467567	467568	467742	467830	467831	467832	467866	468004	468005	468006	468074	469157	469158	469317	469318	469319	469320	469389	469390	469439	469440	470334	470335	470336	470337	470338	470339	470340	470341	470342	470343
BAC	Name	F5K20																																			
BAC	some Length	112929	112929	112929	112929	112929	112929	112929	112929	112929	112929	112929	112929	112929	112929	112929	112929	112929	112929	112929	112929	112929	112929	112929	112929	112929	112929	112929	112929	112929	112929	112929	112929	112929	112929	112929	112929
	Chromosome	33	3	33	3	33	3	3	3	3	co	33	n	33	3	3	3	3	3	3	3	3	33	3	3	3	33	c,	3	3	3	3	3	3	3	33	က
	Seq id	AL132960																																			
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Columbia/	Landsberg	-3/3	-3/3	12/-12	L-/L	3/-3	-1/1	1/-1	-1/1	1/-1	1/-1	1/-1																									
	Method	2	7	2	2	2	-	_	1			-	-	_	_			П	-	1		1	-	1			_	_	_	1	_	_		_	_	_	_
	Type	N N	ONI	ONI	ON N	ONI	IND	ONI	ONI	ONI	ONI	ONI	SNP																								
	Right	68409	68410	729	86093	89296	10430	25599	26007	30831	42929	56574	53825	28614	22292	22169	85756	85655	85058	96225	83733	83929	73060	73183	7482	7651	7631	7616	2097	7618	46545	30615	59340	60143	82703	82826	81719
	Left	68408	68409	716	86085	96764	10429	25597	26006	30829	42927	56572	53823	28612	22290	22167	85754	85653	85056	96223	83731	83927	73058	73181	7480	7649	7629	7614	9092	7616	46543	30613	59338	60141	82701	82824	81717
Marker	Name	470344	470345	470346	470347	470348	471345	471346	471347	471348	471349	471350	466961	466962	467003	467004	467043	467044	467045	467049	467122	467123	467296	467297	467330	467331	467332	467333	467334	467335	467515	467620	468087	468088	468466	468467	468468
BAC	Name	F5K20	F24B22																																		
BAC	Chromosome Length	112929	112929	112929	112929	112929	112929	112929	112929	112929	112929	112929	100285	100285	100285	100285	100285	100285	100285	100285	100285	100285	100285	100285	100285	100285	100285	100285	100285	100285	100285	100285	100285	100285	100285	100285	100285
	Chromos	3	3	3	3	3	3	3	3	3	3	3	3	3	3	3	3	3	3	3	e	3	3	3	3	3	3	3	3	3	3	3	3	3	3	3	3
	Seq id	AL132960	AL132957																																		
Sed	unu	102	102	102	102	102	102	102	102	102	102	102	103	103	103	103	103	103	103	103	103	103	103	103	103	103	103	103	103	103	103	103	103	103	103	103	103

Columbia/	Landsberg	A/G	G/T	C/A	A/C	C/G	G/A	A/G	A/G	C/G	C/T	G/T	CT	A/G	C/T	T/C	A/G	T/C	G/A	C/T	T/C	T/A	A/G	T/A	G/T	CT											
Columbia/	Landsberg																										-3/3	4/-4	9-/9	9-/9	101/-101	9-/9	11/-11	LIL-	6/6-	6/6-	-13/13
	Method	1		_	1	1	1	<del></del>			1	-1	_	1	_	_	-			_		_		-		_	2	2	2	2	2	2	7	2	2	2	2
	Type	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP			ONI	IND	ONI	ON.	ONI	ONI	ONI		ONI						
	Right	83052	82609	19848	19286	19708	3932	3858	3798	3650	4890	4421	4076	58260	58353	56390	57374	16899	93314	93623	64200	70802	96202	86684	86518	71836	1072	34720	509	65594	78202	78895	89427	89481	90003	90006	86606
	Left	83050	82607	19846	19284	19706	3930	3856	3796	3648	4888	4419	4074	58258	58351	56388	57372	16897	93312	93621	64198	70800	70794	86682	86516	71834	1071	34715	502	65587	78100	78888	89415	89480	90005	90005	26606
Marker	Name	468469	468470	468904	468905	468906	468922	468923	468924	468925	468926	468927	468928	468955	468956	468957	468958	468969	469035	469036	469088	469305	469306	469346	469347	469456	469888	469889	469890	469891	469892	469893	469894	469895	469896	469897	469898
BAC	Name	F24B22	F24B22	F24B22	F24B22	F24B22	F24B22	F24B22	F24B22	F24B22	F24B22	F24B22	F24B22	F24B22	F24B22	F24B22	F24B22	F24B22	F24B22	F24B22	F24B22	F24B22	F24B22	F24B22	F24B22	F24B22	F24B22	F24B22	F24B22	F24B22	F24B22						
BAC	osome Length	100285	100285	100285	100285	100285	100285	100285	100285	100285	100285	100285	100285	100285	100285	100285	100285	100285	100285	100285	100285	100285	100285	100285	100285	100285	100285	100285	100285	100285	100285	100285	100285	100285	100285	100285	100285
	Chromosome	3	3	3	33	3	3	3	3	3	3	3	33	$\varepsilon$	3	ж	3	3	3	3	3	3	3	3	3	3	3	3	33	3	3	3	3	3	3	3	es
	Seq id	AL132957	AL132957	AL132957	AL132957	AL132957	AL132957	AL132957	AL132957	AL132957	AL132957	AL132957	AL132957	AL132957	AL132957	AL132957	AL132957	AL132957	AL132957	AL132957	AL132957	AL132957	AL132957	AL132957	AL132957	AL132957	AL132957	AL132957	AL132957	AL132957	AL132957						
Sed	unu	103	103	103	103	103	103	103	103	103	103	103	103	103	103	103	103	103	103	103	103	103	103	103	103	103	103	103	103	103	103	103	103	103	103	103	103

Columbia/	Landsberg															G/A	A/C	A/G	C/T	C/A	T/C	T/A	G/T	G/A	A/G	G/T	G/A	C/A	G/T	A/C	G/A	A/C	C/G	T/G	T/G	A/T	ΑΤ
Columbia/	Landsberg	L/L-	6-/6	-5/5	-1/1	-1/1	-1/1	1/-1	3/-3	2/-2	1/-1	1/-1	2/-2	4/-4	-3/3																						
	Method	2	7		_	-	_		1	1	1	1	1	2	2	1	-	_		<del></del> 4	-	-	1	1	1	1			-		-	_	-	_	_		<b>—</b>
	Type	ONI	IND	ON!	ONI	IND	IND	IND	IND	ONI	IND	ONI	ONI	ONI	IND	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP
	Right	91402	91514	9397	16901	22288	30311	34717	34721	44892	63624	71932	7672	5273	6622	65728	64555	13413	13411	51285	58242	969	1030	9698	8088	74560	52921	52247	75062	11853	83504	83057	83147	82830	82879	82945	82241
	Left	91401	91504	9366	16900	22287	30310	34715	34717	44889	63622	71930	6992	5268	6621	65726	64553	13411	13409	51283	58240	594	1028	8694	9088	74558	52919	52245	75060	11851	83502	83055	83145	82828	82877	82943	82239
Marker	Name	469899	469900	469901	471214	471215	471216	471217	471218	471219	471220	471221	471222	470576	470577	466837	466838	466880	466881	466899	467231	467810	467811	467869	467870	467974	468127	468128	468556	468577	469090	469091	469092	469093	469094	469095	469096
BAC	Name	F24B22	T12E18	T12E18	T15C9	T15C9	T15C9	T15C9	T15C9	T15C9	T15C9	T15C9	T15C9	T15C9	T15C9	T15C9	T15C9	T15C9	T15C9	T15C9	T15C9	T15C9	T15C9	T15C9	T15C9	T15C9											
BAC	Chromosome Length	100285	100285	100285	100285	100285	100285	100285	100285	100285	100285	100285	100285	40766	40766	84233	84233	84233	84233	84233	84233	84233	84233	84233	84233	84233	84233	84233	84233	84233	84233	84233	84233	84233	84233	84233	84233
	Chrom	3	3	3	3	3	$\alpha$	3	3	3	3	3	$\mathcal{E}$	3	3	$\alpha$	3	$\varepsilon$	3	3	$\mathfrak{S}$	$\varepsilon$	$\alpha$	$\alpha$	3	3	$\varepsilon$	3	3	3	33	3	3	3	3	3	e
	Seq id	AL132957	AL132971	AL132971	AL132970	AL132970	AL132970	AL132970	AL132970	AL132970	AL132970	AL132970	AL132970	AL132970	AL132970	AL132970	AL132970	AL132970	AL132970	AL132970	AL132970	AL132970	AL132970	AL132970	AL132970	AL132970											
Sed	unu	103	103	103	103	103	103	103	103	103	103	103	103	104	104	105	105	105	105	105	105	105	105	105	105	105	105	105	105	105	105	105	105	105	105	105	105

SNP Base Columbia/ Landsberg G/T C/T C/T A/G A/T T/A C/G C/G C/G A/T T/A A/T T/A A/T T/A A/T T/A A/T T/A A/G A/T T/A T/A
Indel Size Columbia/ Landsberg 4/-4 -4/4 9/-9 9/-9 9/-9 18/-18 -3/3 11/-11 11/-11 -28/28 4/-4 -1/1
Method
Right 82619 82657 82878 20703 23055 38785 39774 40007 42786 42746 68817 73030 81444 50545 72882 15268 14651 14672 1960 19761 19761 19760 19778 9332 8287 4027
Left 82617 82655 82876 20698 23054 38775 39764 40002 42745 42888 47779 50544 72880 15744 14670 15266 15744 19658 19759 19759 19759 19759 19759 19759 19759 19776 9330
Marker Name 469097 469098 469099 470600 470601 470602 470603 470603 470604 470609 470609 470609 470609 470610 470610 470610 470610 470610 470610 470610 470610 470610 470610 470610 470610 470610 470610 470610 470610 470610 466804 466806 466806 466806 466806 466806 466806 466806 466806 466806 467445 467448 467448 467448 467449 467448 467449 467429 468557
BAC Name T15C9 T15C112 T26I12
Chromosome Length 3 3 44233 44233 3 44233 44233 44233 44233 44233 44233 44233 44233 44233 44233 44233
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Seq id ALI32970 ALI32970 ALI32970 ALI32970 ALI32970 ALI32970 ALI32970 ALI32970 ALI32970 ALI32970 ALI32970 ALI32970 ALI32954
Seq 105 105 105 106 106 106 106 106 106 106 106 106 106

Columbia/	Landsberg	T/C	C/G	T/G	C/T	G/A	C/T	T/C	T/C	C/A	T/A	T/C	G/C																				C/A	C/A	G/C	A/C	C/G
Columbia/	Landsberg														9/9-	-11/11	-14/14	388/-388	16/-16	-3/3	16/-16	9-/9	4/-4	3/-3	5/-5	09/09-	-1/1	4/-4	3/-3	-1/1	-1/1	-1/1					
	Method	-	T	1	1	1	-	_	1		_		_	2	2	2	2	2	2	2	2	2	2	2	2	2	-	-	1	1	_		<del></del>	-	1	1	
	Type	SNP	IND		IND	ONI		IND	ONI	ONI	ONI	QNI	ONI	IND	ONI	QNI	ON!	ONI	ONI	ONI	ONI	SNP	SNP	SNP	SNP	SNP											
	Right	4249	4785	4448	4225	10028	10115	7367	7275	14014	13747	13848	14158	11420	22608	23944	33858	35329	40031	43067	44213	46962	4726	4890	63286	69/99	27636	4732	4895	62260	62209	87498	5087	5347	5155	5003	5158
	Left	4247	4783	4446	4223	10026	10113	7365	7273	14012	13745	13846	14156	11413	22607	23943	33857	34940	40014	43066	44196	46955	4721	4886	63280	89/99	27635	4727	4891	62259	80/59	87497	5085	5345	5153	5001	5156
Marker	Name	468559	468560	468561	468562	468939	468940	468979	468980	469010	469011	469012	469013	470874	470875	470876	470877	470878	470879	470880	470881	470882	470883	470884	470885	470886	471555	471556	471557	471558	471559	471560	466802	466803	466804	466805	466806
BAC	Name	T26112	T26112	T26112	T26112	T26112	T26I12	T26I12	T26112	T26112	T26I12	T26112	T26112	T26112	T26112	T26112	T26112	T26I12	T26112	T26112	T26112	T26I12	T26112	T26112	T26112	T22E16	T22E16	T22E16	T22E16	T22E16							
BAC	Chromosome Length	26688	26688	26688	288997	26688	26688	26688	26688	26688	26688	26688	26688	26688	26688	26688	26688	88997	26688	26688	26688	26688	26688	26688	26688	26688	26688	26688	26688	26688	26688	26688	103240	103240	103240	103240	103240
	ū	54 3	3			3			3	54	3	3	3	3	3	54	3		5.4		. <del>4</del> 3			3	3	4.3	3	3	3		3	5.4	5 3		5 3	5 3	5 3
	Seq id	AL132954	AL132954	AL132954	AL132954	AL132954	AL132954	AL132954	AL132954	AL132954	AL132954	AL132954	AL132954	AL132975	AL132975	AL132975	AL132975	AL132975																			
Seq	unu	106	901	106	106	106	106	106	106	106	106	901	106	106	106	106	106	106	106	106	106	106	106	106	106	106	106	106	106	106	106	106	107	107	107	107	107

SNP Base Columbia/	Landsberg	J/C	A/G	CT	T/A	T/C	A/C	T/C	T/C	A/G	A/G	$C/\Gamma$	T/A	C/A	G/A	T/C	G/A	T/C	A/C	C/G	C/T	CT	A/G	CT	G/A	G/A	T/A	T/C	A/T	A/T	T/A	T/A	T/C	CT	C/G	C/T	C/A
Indel Size Columbia/	Landsberg																																				
	Method	_	<b></b>	_	<b>.</b> —1	1	1	1	_	_		-	1	1	_						1	<b>⊢</b> ⊣	<b>—</b>	1	_		1	1				_	_	1	1	-	
	Type	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP
	Right	5127	5119	89248	21900	22006	21994	21985	21974	21750	22034	21909	33037	32559	11078	27966	100027	100899	101620	20666	101100	27401	81866	65703	30286	30324	30577	30548	30388	30389	98246	99051	98763	98144	53654	85894	37305
	Left	5125	5117	89246	21898	22004	21992	21983	21972	21748	22032	21907	33035	32557	11076	27964	100025	100897	101618	99905	101098	27399	81864	65701	30284	30322	30575	30546	30386	30387	98244	99049	98761	98142	53652	85892	37303
Marker	Name	466807	466808	466932	467013	467014	467015	467016	467017	467018	467019	467020	467155	467156	467227	467355	467413	467414	467415	467416	467417	467585	467820	467858	468079	468080	468081	468082	468083	468084	468318	468319	468320	468321	468381	468382	468448
BAC	Name	T22E16	T22E16	T22E16	T22E16	T22E16	T22E16	T22E16	T22E16	T22E16	T22E16	T22E16	T22E16	T22E16	T22E16	T22E16	T22E16	T22E16	T22E16	T22E16	T22E16	T22E16	T22E16	T22E16	T22E16	T22E16	T22E16	T22E16	T22E16	T22E16	T22E16	T22E16	T22E16	T22E16	T22E16	T22E16	T22E16
BAC	Chromosome Length	103240	103240	103240	103240	103240	103240	103240	103240	103240	103240	103240	103240	103240	103240	103240	103240	103240	103240	103240	103240	103240	103240	103240	103240	103240	103240	103240	103240	103240	103240	103240	103240	103240	103240	103240	103240
	Chron	$\alpha$	n	n	8	$\varepsilon$	$\epsilon$	c	3	3	$\omega$	Э	3	ю	Э	33	Э	$\epsilon$	3	3	3	3	3	3	3	3	3	$\mathfrak{C}$	3	3	3	3	3	3	3	$\varepsilon$	3
	Seq id	AL132975	AL132975	AL132975	AL132975	AL132975	AL132975	AL132975	AL132975	AL132975	AL132975	AL132975	AL132975	AL132975	AL132975	AL132975	AL132975	AL132975	AL132975	AL132975	AL132975	AL132975	AL132975	AL132975	AL132975	AL132975	AL132975	AL132975	AL132975	AL132975	AL132975	AL132975	AL132975	AL132975	AL132975	AL132975	AL132975
Sea	unu	107	107	107	107	107	107	107	107	107	107	107	107	107	107	107	107	107	107	107	107	107	107	107	107	107	107	107	107	107	107	107	107	107	107	107	107

Columbia/	Landsberg	G/A	T/C	A/G	A/C	T/C	A/G	T/A	A/G	G/A	C/A	G/T	G/T																								
Columbia/	Landsberg													12/-12	3/-3	21/-21	12/-12	6-/6	-64/64	9-/9	-16/16	13/-13	L-1L	9/9-	9/9-	22/-22	-10/10	-10/10	8-/8	3/-3	3/-3	4/-4	-17/17	-3/3	3/-3	1/-1	-1/1
	Method	1	-	_			_	-	_	_	_	-	<b>-</b>	2	2	2	2	2	2	2	2	2	2	2	2	2	2	2	2	2	2	2	2	2		1	<b>-</b>
	Type	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	ON.	ONI	ON.	IND	ONI	ONI	IND	ONI	ONI	ONI	ONI	ONI	ON N	ON			ONI		ONI	ONI		ONI	ON	ON O
	Right	33614	50679	50736	14986	83504	83604	43887	44087	3288	3484	3803	51519	102522	15419	23206	31098	39643	42042	43599	53705	57497	60208	60209	60300	60885	60961	60963	61422	64586	72346	74821	93775	08866	15425	30357	33721
	Left	33612	20677	50734	14984	83502	83602	43885	44085	3286	3482	3801	51517	102509	15415	23184	31085	39633	42041	43592	53704	57483	60200	60298	60299	60862	09609	60962	61413	64582	72342	74816	93774	62866	15421	30355	33720
Marker	Name	468706	468810	468811	469086	469311	469312	469321	469322	469393	469394	469395	469457	470793	470794	470795	470796	470797	470798	470799	470800	470801	470802	470803	470804	470805	470806	470807	470808	470809	470810	470811	470812	470813	471518	471519	471520
BAC	Name	T22E16	T22E16	T22E16	T22E16	T22E16	T22E16	T22E16	T22E16	T22E16	T22E16	T22E16	T22E16	T22E16	T22E16	T22E16	T22E16	T22E16	T22E16	T22E16	T22E16	T22E16	T22E16	T22E16	T22E16	T22E16	T22E16	T22E16	T22E16	T22E16	T22E16	T22E16	T22E16	T22E16	T22E16	T22E16	T22E16
BAC	Chromosome Length	103240	103240	103240	103240	103240	103240	103240	103240	103240	103240	103240	103240	103240	103240	103240	103240	103240	103240	103240	103240	103240	103240	103240	103240	103240	103240	103240	103240	103240	103240	103240	103240	103240	103240	103240	103240
	ט			33					33		3	3	c					3				3			3	3									3		
	Seq id	AL 132975	AL132975																																		
Sed	unu	107	107	107	107	107	107	107	107	107	107	107	107	107	107	107	107	107	107	107	107	107	107	107	107	107	107	107	107	107	107	107	107	107	107	107	107

Columbia/	Landsberg				T/A	G/C	C/A	G/A	C/A	T/G	A/G	T/G	A/G	A/T	G/T	CT	A/G	T/A	A/G	T/A	A/C	T/C	C/T	T/A	A/C	A/C	C/T	CT	C/A	T/C	C/G	A/G	C/G	Α⁄Τ	G/T	G/T	C/T
Columbia/	Landsberg	1/-1	-1/1	-3/3																																	
	Method	1	1	1	<b>.</b>	1	1	1	_		_	1	_	<del></del>	1	-				_	1		1	_	_	1	1	_	1	1	_	1	-	_	1	<del></del>	1
	Type	QNI	ONI	ONI	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP									
	Right	53597	66286	99881	84227	84085	63738	63880	64342	64369	64371	64432	64433	63963	64188	64274	34477	33765	55883	56892	29695	26898	56958	67612	66116	67156	66092	67224	13256	12381	11999	13570	13144	12232	12032	13602	13262
	Left	53595	86286	08866	84225	84083	63736	63878	64340	64367	64369	64430	64431	63961	64186	64272	34475	33763	55881	26890	56963	96895	56956	67610	66114	67154	06099	67222	13254	12379	11997	13568	13142	12230	12030	13600	13260
Marker	Name	471521	471522	471523	472053	472054	472076	472077	472078	472079	472080	472081	472082	472083	472084	472085	472188	472189	472202	472203	472204	472205	472206	472620	472621	472622	472623	472624	472644	472645	472646	472647	472648	472649	472650	472651	472652
BAC	Name	T22E16	T22E16	T22E16	F2809	F2809	F2809	F28O9	F2809	F2809	F28O9	F28O9	F28O9	F2809	F2809	F28O9	F28O9	F28O9	F28O9	F28O9	F28O9	F2809	F28O9	F2809	F28O9	F28O9	F2809	F28O9	F28O9	F28O9	F2809						
BAC	Chromosome Length		3 103240	3 103240	3 85710	3 85710	3 85710	3 85710	3 85710	3 85710	3 85710	3 85710	3 85710	3 85710	3 85710	3 85710	3 85710	3 85710	3 85710	3 85710	3 85710	3 85710	3 85710	3 85710	3 85710	3 85710	3 85710	3 85710	3 85710	3 85710	3 85710	3 85710	3 85710	3 85710	3 85710	3 85710	3 85710
	Seq id	975			AL137080	AL137080	AL137080	AL137080	AL137080	AL137080		AL137080		AL137080	AL137080																						
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Columbia/	Landsberg	C/A	C/A	T/C	A/T	T/C	T/C	T/G	A/T	C/A	T/C	G/C	C/T	A/G	T/A	C/T	C/I	C/G	A/G	T/G	C/A	C/A	A/C	T/G	C/T	G/T	G/T	C/T									
Columbia/	Landsberg																											;	-3/3	476/-476	L-1L	-12/12	9/9-	4/-4	9/9-	4/-4	3/-3
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	Right	47342	51450	73345	73613	11556	48966	48216	48198	3348	2891	2290	4693	78084	82153	81751	17777	68828	89889	68819	69583	20007	69591	69585	69663	69669	70017	69593	23229	26078	58130	58263	67582	69250	78604	79833	13261
	Left	47340	51448	73343	73611	11554	48964	48214	48196	3346	2889	2288	4691	78082	82151	81749	17775	68826	99889	68817	69581	90002	68569	69583	69661	<i>L</i> 9669	70015	69591	23228	25601	58122	58262	67581	69245	78603	79828	13257
Marker	Name	472763	472869	472927	472928	473117	473281	473282	473283	473436	473437	473438	473439	473512	473613	473614	473761	473777	473778	473779	473780	473781	473782	473783	473784	473785	473786	473787	473891	473892	473893	473894	473895	473896	473897	473898	474202
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BAC	Chromosome Length	85710	85710	85710	85710	85710	85710	85710	85710	85710	85710	85710	85710	85710	85710	85710	85710	85710	85710	85710	85710	85710	85710	85710	85710	85710	85710	85710	85710	85710	85710	85710	85710	85710	85710	85710	85710
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	Seq id	AL137080	AL137080	AL137080	AL137080	AL137080	AL137080	AL137080	AL137080	AL137080	AL137080	AL137080	AL137080																								
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SNP Base Columbia/ Landsberg	A/G C/T 17/C 17/G 17/G A/T C/A A/G A/C A/C A/C A/C A/C C/T C/A
Indel Size Columbia/ Landsberg 2/-2 1/-1 -2/2 1/-1 1/-1 1/-1 1/-1 -1/1 -1/	
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Right 13265 13266 17449 34188 42974 48976 56924 56931 57011 67585 67586 69696	24229 54790 56630 56767 16944 16904 16906 23458 23343 24330 24684 77123 10336 15202 68149 45990 64254
Left 13262 13264 17448 34186 42972 48448 48974 56923 56929 57009 67585 69694 73585	2428 5628 56628 56765 16942 16902 16904 23456 23456 23456 2341 2482 77121 10334 15200 68147 45988
Marker Name 474203 474204 474206 474206 474200 474210 474211 474213 474213 474213 474213	4/4219 4/6886 4/6888 4/67105 4/67106 4/67109 4/67111 4/67111 4/67111 4/67111 4/67111 4/67111 4/67111 4/67113 4
BAC Name F2809 F2809 F2809 F2809 F2809 F2809 F2809 F2809 F2809 F2809 F2809 F2809 F2809 F2809	7.88.09 7.88.10 7.88.110 7.88.110 7.88.110 7.88.110 7.88.110 7.88.110 7.88.110 7.88.110 7.88.110 7.88.110 7.88.110 7.88.110 7.88.110
BAC Chromosome Length 3 85710 3 85710	87.10 87503 87503 87503 87503 87503 87503 87503 87503 87503 87503 87503 87503 87503 87503 87503 87503 87503
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Indel Size Columbia/ Landsberg -4/4 -7/7 -4/4 3/-3 10/-10 -4/4 -9/9 -4/4 -6/6 11/-11 -2/2 1/-1 -1/1 -1/1	i
Method	(
	SNP SNP SNP SNP SNP SNP SNP
Right 13776 59426 59327 85768 85736 85736 85678 30101 30094 49989 4983 228159 30319 55861 56788 57267 59518 66315 73952 75810 16900 29967 40487 40487 56786 56786 56786	108184 32788 32636 32968 32741 32886 60144
Left 13774 59424 59325 85766 85734 85505 85676 30099 30092 49987 49831 28158 30318 55860 56784 57256 59517 63163 66314 73951 75798 16899 29965 40488 56784	108182 32786 32634 32966 32739 32884 60142
Marker Name 468261 468714 468715 469336 469337 469339 469403 469403 469403 469403 471032 471033 471035 471036 471037 471039 471039 471040 471615 471615 471615	467534 468299 468300 468301 468302 468303 468735
BAC Name 18H10 18H10 18H10 18H10 18H10 18H10 18H10 18H10 18H10 18H10 18H10 18H10 18H10 18H10 18H10 18H10 18H10 18H10 18H10	T10K17 T10K17 T10K17 T10K17 T10K17 T10K17
BAC some Length 87503	109016 109016 109016 109016 109016 109016
Chromosome	
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SNP Base Columbia/ Landsberg A/T	A/C 1/C A/T 1/A 1/C A/G	T/C G/A G/T	G/A G/A C/A A/T C/T C/G G/A	A/C A/C C/T 1/C 1/C 1/C 6/C
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Right 102182 19003 19018 18783 31428 39595	11347 66980 22555 79392 33181	57131 59852 25322 23229 23550	10283 9795 11114 10590 9842 10970 11276 30261 30490 60779	61793 61792 61568 50454 12280 12382 32370 32492
Left 102180 17453 17468 18775 31427 39594	11345 66978 22553 79390 33179	57129 59850 25320 23227 23548	10281 9793 11112 10588 9840 10968 11274 30259 30488	61791 61790 61566 50452 12278 12380 32368 32490
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SNP Base Columbia/ Landsberg G/C G/A C/G C/A T/A T/A G/C C/G C/G C/T T/A T/A T/A T/A T/A T/A T/A T/A T/A T	C/T G/A A/G A/G C/A G/C
Indel Size Columbia/ Landsberg Landsberg -5/5 -10/10 3/-3 -7/7 -1/1 -2/2 -6/6 1/-1 -1/1 -1/1 1/-1 1/-1	
Method 1	
	SNP SNP SNP SNP SNP SNP
Right 31621 1442 1556 43826 43826 43826 43648 36751 66805 66293 64918 65538 62594 59043 58949 123 2394 351 4122 11280 25508 25546 25508 65508	44656 91106 91398 71288 26256 21191 21164 21145
Left 31619 1440 1554 43824 43824 43646 36806 36749 66291 64916 65536 62592 59041 122 2393 347 4121 11222 11279 25544 25611 44217 65506	24654 91104 91396 71286 26254 21189 21162 21143
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BAC Name T16L24	T20K12 T20K12 T20K12 T20K12 T20K12 T20K12 T20K12 T20K12
BAC Chromosome Length 3 91851 91851 3 91851 3 91851 3 91851 3 91851 3 91851 3 91851 3 91851 3 91851 3 91851 3 91851 3 91851 3 91851 3 91851 3 91851 3 91851 3	109155 109155 109155 109155 109155 109155 109155 109155
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Columbia/	Landsberg	C/T	A/C	AT	A/C	T/C	A/G	A/G	C/T	A/C	A/C	A/T	T/A	G/T	C/T	C/T	G/A	A/C	T/C	T/C	G/A	A/C	A/G	A/C	A/T	T/C	A/C	TVC	C/G	C/T	A/T	A/T	A/T	T/A	T/C	و <i>ر</i> د دور	A/G
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	Left	21175	90221	81292	13315	13295	13576	13329	13537	65821	36927	37186	35691	36429	35961	36486	72533	72046	72199	72393	67307	90829	67242	64427	64428	29387	29270	29545	29243	30245	69139	69138	23110	76364	76231	76360	77568
Marker	Name	472234	472346	472376	472536	472537	472538	472539	472540	472678	472727	472728	472729	472730	472731	472732	472897	472898	472899	472900	473025	473026	473027	473387	473388	473428	473429	473430	473431	473432	473451	473452	473505	473607	473608	473609	473610
BAC	Name	T20K12	T20K12	T20K12	T20K12	T20K12	T20K12	T20K12	T20K12	T20K12	T20K12	T20K12	T20K12	T20K12	T20K12	T20K12	T20K12	T20K12	T20K12	T20K12	T20K12	T20K12	T20K12	T20K12	T20K12	T20K12	T20K12	T20K12	T20K12	T20K12	T20K12	T20K12	T20K12	T20K12	T20K12	T20K12	T20K12
BAC	Chromosome Length			3 109155	3 109155	3 109155	3 109155	3 109155	3 109155	3 109155	3 109155	3 109155	3 109155	3 109155	3 109155	3 109155	3 109155	3 109155	3 109155	3 109155	3 109155	3 109155	3 109155	3 109155	3 109155	3 109155	3 109155	3 109155	3 109155	3 109155	3 109155	3 109155	3 109155	3 109155		3 109155	3 109155
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SNP Base Columbia/ Landsberg A/T C/T A/C C/T C/T G/A G/A	17.A 17.C 17.C 17.A 17.A 17.A 17.A 17.A 17.A 17.A 17.A
Indel Size Columbia/ Landsberg -9/9 24/-24 1/-1 1/-1 1/-1 -1/1 -1/1 -1/1 -1/1 -1/	<b>1</b> 77-
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Right 76330 76330 76390 94155 94135 33570 24472 24549 13778 13778 13778 13780 24494 26419 29645 36102 52680 68864 69144	34221 35345 35246 34484 35377 30407 30572 80166 10920 10937 88669 23931
Left 76328 76388 94153 94153 94133 33568 24470 24547 52382 2227 2301 13776 13776 13778 24492 24492 26418 29644 36101 52679 68863 69142	34219 34219 35343 35244 34482 35375 30405 30570 80164 10918 10935 88667
Marker Name 473611 473612 473618 473619 473619 473740 473747 474421 474423 474424 474429 474429 474429 474431 474433	4/4434 466792 466793 466794 466795 466955 466956 467349 467395 467395
BAC Name T20K12	120K12 F2A19 F2A19 F2A19 F2A19 F2A19 F2A19 F2A19 F2A19
BAC Chromosome Length 109155 3 109155 3 109155	109155 95993 95993 95993 95993 95993 95993 95993 95993 95993
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SNP Base Columbia/	Landsberg	C.C.	T/A	A/I	A/T	A/T	C/G	G/A	T/G	G/T	G/T	A/T	G/T	G/A	A/C	T/C	G/C	A/T	AVT	A/T	T/A	G/A	T/A	T/G	A/G	C/T	A/I	A/T	G/I	G/A	G/A	C/A	T/C	T/C	A/G	C/G	<u>5</u> /2
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	Right	42660	13719	13702	13737	13743	23141	43685	43641	43572	49656	38747	38755	72110	71951	72678	72472	72731	72668	72667	19794	19645	19661	19608	20052	19804	19809	19660	26560	80838	90564	90017	89806	2268	90871	90757	90734
	Left	42658	13717	13700	13735	13741	23139	43683	43639	43570	49654	38745	38753	72108	71949	72676	72470	72729	72666	72665	19792	19643	19659	19606	20050	19802	19807	19658	56558	90836	90562	90015	99806	89975	69806	90755	90732
Marker	Name	467681	467795	467796	467797	467798	468092	468114	468115	468116	468359	468396	468397	468544	468545	468546	468547	468548	468549	468550	468687	468688	468689	468690	468691	468692	468693	468694	468708	468824	468825	468826	468827	468828	468829	468830	468831
BAC	Name	F2A19	F2A19	F2A19	F2A19	F2A19	F2A19	F2A19	F2A19	F2A19	F2A19	F2A19	F2A19	F2A19	F2A19	F2A19	F2A19	F2A19	F2A19	F2A19	F2A19	F2A19	F2A19	F2A19	F2A19	F2A19	F2A19	F2A19	F2A19	F2A19	F2A19	F2A19	F2A19	F2A19	F2A19	F2A19	F2A19
BAC	Chromosome Length	95993	95993	95993	95993	95993	95993	95993	95993	95993	95993	95993	95993	95993	95993	95993	95993	95993	95993	95993	95993	95993	95993	95993	95993	95993	95993	95993	95993	95993	95993	95993	95993	95993	95993	95993	95993
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	Seq id	AL 132962	AL132962	AL132962	AL132962	AL132962	AL132962	AL132962	AL132962	AL132962	AL132962	AL132962	AL132962	AL132962	AL132962	AL132962	AL132962	AL132962	AL132962	AL132962	AL132962	AL132962	AL132962	AL 132962	AL132962	AL132962	AL132962	AL132962	AL132962	AL132962	AL132962	AL132962	AL132962	AL132962	AL132962	AL132962	AL132962
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SNP Base Columbia/ Landsberg A/T A/T C/T C/T C/T T/C C/T T/C C/T T/C
Indel Size Columbia/ Landsberg 4/4 3/-3 -7/7 14/-14 -13/13 33/-3 -9/9 4/-4 -3/3 -1/1 1/-1 -2/2 -1/1 2/-2 2/-2 2/-2 2/-2
Method
Right 90607 90557 89956 4648 4665 75111 74725 30987 15446 10976 13306 17592 26456 32218 37330 41146 43530 47705 52149 58490 65624 79230 89713 17594 17598 19593 20540 38554 38571 43533
Left 90605 90555 89954 4646 4663 75109 74723 30985 15444 10975 13302 17591 26441 32217 37319 37576 41138 43523 47701 52148 58489 65619 79229 89712 17597 17597 17597 17593 38567 42508 43530
Marker Name 468832 468833 468833 468833 468833 469833 469076 469076 470109 470109 470109 470111 470111 470111 470111 470118 471283 471289 471289 471290 471291 471299
BAC Name F2A19
BAC Chromosome Length 3 3 4 55993 3 959993 3 95993 3
OF COLUMN SOURCE
Seq id AL132962
Seq num

SNP Base Columbia/	Landsberg		į	I/C	G/C	A/G	A/G	C/G	C/A	T/C	T/C	C/A	C/A	G/C														į	A/C	G/A	T/C	G/C	A/G	A/G	C/T	C/T	G/T
Indel Size Columbia/	Landsberg	1/-1	2/-2												12/-12	-5/5	-3/3	18/-18	18/-18	186/-186	4/-4	19/-19	3/-3	9/9-	9/9-	1/-1	-1/1	-1/1									
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	Right	74950	80227	18924	70639	71012	60688	86419	74320	83380	83921	17705	17704	17822	21131	68731	6932	71602	71627	77369	77903	78410	81281	82207	82209	23410	90773	92366	43053	69557	69553	69951	69737	69881	69954	70033	70112
	Left	74948	80224	18922	70637	71010	88907	86417	74318	83378	83919	17703	17702	17820	21118	68730	6931	71583	71608	77182	77898	78390	81277	82206	82208	23408	90772	97365	43051	69555	69551	69649	69735	62869	69952	70031	70110
Marker	Name	471295	471296	466925	466979	466980	467350	468252	468663	468767	468768	468862	468863	468864	469675	469676	469677	469678	469679	469680	469681	469682	469683	469684	469685	471148	471149	471150	471740	471898	471899	471900	471901	471902	471903	471904	471905
BAC	Name	F2A19	F2A19	F15G16	F15G16	F15G16	F15G16	F15G16	F15G16	F15G16	F15G16	F15G16	F15G16	F15G16	F15G16	F15G16	F15G16	F15G16	F15G16	F15G16	F15G16	F15G16	F15G16	F15G16	F15G16	F15G16	F15G16	F15G16	F25E4								
BAC	Chromosome Length	95993	95993	104868	104868	104868	104868	104868	104868	104868	104868	104868	104868	104868	104868	104868	104868	104868	104868	104868	104868	104868	104868	104868	104868	104868	104868	104868	80019	80019	80019	80019	80019	80019	80019	80019	80019
	Chro	$\alpha$	$\epsilon$	$\epsilon$	$\varepsilon$	$\epsilon$	3	3	3	3	3	3	3	$\mathcal{E}$	$\epsilon$	3	$\epsilon$	3	$\varepsilon$	ĸ	ري ا	$\varepsilon$	3	$\epsilon$	$\kappa$	т	$\kappa$	$\epsilon$	4	4	4	4	4	4	4	4	4
	Seq id	AL132962	AL132962	AL132959	AL132959	AL132959	AL132959	AL132959	AL132959	AL132959	AL132959	AL132959	AL132959	AL132959	AL132959	AL132959	AL132959	AL132959	AL132959	AL132959	AL132959	AL132959	AL132959	AL132959	AL132959	AL132959	AL132959	AL132959	AL050399								
Seq	unu	115	115	116	116	116	116	116	116	116	116	116	116	116	116	116	116	116	116	116	116	116	116	116	116	116	116	116	117	117	117	117	117	117	117	117	117

SNP Base Columbia/ Landsberg T/A C/A T/C A/G A/G A/T T/A T/A T/A T/A T/A T/A T/A T/A T/A	A/T A/T A/T T/C C/T
Indel Size Columbia/ Landsberg  -1/1 1/-1 -2/2 -1/1	c/c-
Method	
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Right 41868 41615 42206 41642 41642 41609 41675 31025 31026 31027 31028 31029 32571 27340 27437 27644 27645 27645 27645 27645	2/64/ 24904 10082 10084 82859 82786
Left 41866 41613 42204 41640 41640 41640 41673 32245 31024 31023 31025 31026 31027 22948 27338 27435 27641 27644 27644 27644 27644 27644 27644 27644 27644 27644	27646 24902 10080 10082 82857 82784
Marker Name 472029 472030 472031 472031 472033 472033 472033 472033 472034 472034 472175 472177 472179 472179 473090 473090 473099 473099 473099 473099 473096 473096 473096 473096 473096 473099	474160 471656 471709 471710 471718
BAC Name F25E4	F25E4 F718 F718 F718 F718 F718
BAC Chromosome Length 80019 4 80019 4 80019	80019 114759 114759 114759 114759
O 4 4 4 4 4 4 4 4 4 4 4 4 4 4 4 4 4 4 4	4 v v v v v
Seq id ALO50399	AL050399 AL137189 AL137189 AL137189 AL137189
Seq 1117 1117 1117 1117 1117 1117 1117 11	117 118 118 118 118

SNP Base Columbia/ Landsberg G/C T/C A/G C/T C/T T/C	C/A A/G A/G C/G C/T	7.4 7.7 7.7 7.4 7.4 7.7	C/A 17/C C/G G/A G/A G/A A/C
Indel Size Columbia/ Landsberg			
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Right 11424 91015 91137 90939 90803 48327	47560 47618 21006 21117 81468 52759 52904 89865 90193	101611 16716 14990 14796 55681 63852 64013 96197 103954	103712 104045 103712 103872 103592 89249 88844 88802 89303 89245
Left 11422 91013 91135 90937 90907 90801 48325	47558 47616 21004 21115 81466 52757 52902 89863 90191	101609 16714 14988 14794 55679 63850 64011 96195 103952	104043 104043 103710 103870 103590 89247 88842 88800 89301 89301
Marker Name 471754 471760 471761 471762 471763 471764	471947 471948 472042 472157 472157 472218 472228 472229	472272 472479 472480 472481 472662 472711 472712 473100 473253	473254 473255 473256 473257 473260 473261 473262 473264 473265
BAC Name F718 F718 F718 F718 F718 F718 F718	F718 F718 F718 F718 F718 F718 F718	F7.18 F7.18 F7.18 F7.18 F7.18 F7.18 F7.18	F738 F738 F738 F738 F738 F738 F738 F738
BAC Chromosome Length 5 114759 5 114759 5 114759 5 114759 5 114759 5 114759	114759 114759 114759 114759 114759 114759 114759	114759 114759 114759 114759 114759 114759 114759 114759	114759 114759 114759 114759 114759 114759 114759 114759
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Seq 1118 1118 1118 1118 1118	118	118 118 118 118 118	118 118 118 118 118 118 118 118

SNP Base Columbia/ Landsberg G/A G/T C/G	G/T T/A T/C T/C A/G C/A A/G C/G C/G C/T T/A T/A T/A
Indel Size Columbia/ Landsberg -1/1 -1/1 1/-1 1/-1 1/-1 -1/1 -1/1 -1/	7/7-
Method 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1	
	SNP SNP SNP SNP SNP SNP SNP SNP SNP SNP
Right 23892 84568 49975 104026 21301 80668 841 88916 89466 89468 90827 91376 99287	99302 84592 67072 68505 68205 68487 68318 68506 78520 78405 78405 78734 78734 78734 78734 78734 77181 77181
Left 23890 84566 49973 104025 21300 80666 839 88914 89467 90825 91374 96727	99301 84590 67070 68503 68203 68203 68204 78518 78403 78434 78732 78732 78731 77179 771195 771165
Marker Name 473274 473349 473453 474280 474281 474282 474283 474283 474285 474288	474290 472436 473139 473140 473141 473142 473143 473379 473379 473380 473382 473383 473383 473383 473384 473385 473661 473664 473663
BAC Name F718 F718 F718 F718 F718 F718 F718 F718	F7J8 F13G24
BAC Chromosome Length 5 114759 5 114759	88095 88095 88095 88095 88095 88095 88095 88095 88095 88095 88095 88095 88095 88095 88095 88095 88095 88095
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Seq. 1118 1118 1118 1118 1118 1118 1118 11	118 119 119 119 119 119 119 119 119 119

Columbia/	Landsberg	C/C	A/G	S/O	A/G	D/L	C/T													A/G	A/G	A/G	A/G	T/G	A/G	C/T	G/T	C/A	G/A	G/A	A/G	C/A	C/T	T/G			
Columbia/	Landsberg								-3/3	-3/3	-3/3	-15/15	2532/-2532	9-/9	15/-15	-1/1	1/-1	1/-1	2/-2																1/-1	1/-1	1/-1
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	Right	77247	77166	77238	77493	76224	77219	77240	1161	1162	1163	15262	18367	2809	8801	67284	77201	78408	78412	86402	88770	88915	88628	87993	87994	88041	88050	75172	35409	34929	28270	52672	59069	74035	28103	29437	29482
	Left	77245	77164	77236	77491	76222	77217	77238	1160	1161	1162	15261	15834	2802	8785	67283	77199	78406	78409	86400	88768	88913	98628	87991	87992	88039	88048	75170	35407	34927	28268	52670	29062	74033	28101	29435	29480
Marker	Name	473666	473667	473668	473669	473670	473671	473672	473814	473815	473816	473817	473818	473819	473820	474043	474044	474045	474046	471792	471793	471794	471921	471922	471923	471924	471925	472086	472702	472703	473161	473350	473509	473752	474371	474372	474373
BAC	Name	F13G24	F13G24	F13G24	F13G24	F13G24	F13G24	F13G24	T14C9	T14C9	T14C9	T14C9	T14C9																								
BAC	Chromosome Length	88095	88095	88095	88095	88095	88095	88095	88095	88095	88095	88095	88095	88095	88095	88095	88095	88095	88095	110684	110684	110684	110684	110684	110684	110684	110684	110684	110684	110684	110684	110684	110684	110684	110684	110684	110684
	Ch	5	5	5	5	5	5	5	5	5	5	5	S	5	5	5	5	3	5	S	2	5	5	2	5	S	S	2	5	3	3	2	5	5	S	5	2
	Seq id	AL133421	AL133421	AL133421	AL133421	AL133421	AL133421	AL133421	AC006601	AC006601	AC006601	AC006601	AC006601																								
S. C.		119	119	119	119	119	119	119	119	119	119	119	119	119	119	119	119	119	119	120	120	120	120	120	120	120	120	120	120	120	120	120	120	120	120	120	120

	Landsberg Landsberg 1/-1	T/C	A/T	C/T	S/O	C/T	C/T		-11/11	19/-19	4/-4	4/-4	4/-4	9-/9	12/-12	4758/-4758	-17/17	12/-12	-3/3	-3/3	21/-21	5/-5	25/-25	-14/14	-14/14	-2/2		G/A	G/A	G/A	G/A	A/G	A/G	C/T	C/A	C/A
,	Method 1	1	_	T	T	_	1	1	2	2	2	2	2	2	2	2	2	2				2		2	2	<b>-</b>	2	_	_		<b>—</b>	_		<b>-</b>	_	<b></b>
į	Type IND	SNP	ONI	IND	ONI ONI	ONI	ONI	ONI	ONI	ONI	ONI	IND	ONI	IND	ONI	ONI	IND	ON ON	IND	ONI	IND	SNP														
	Right 86765	87042	85826	86846	110733	110765	110925	84898	101611	102611	13924	14468	16221	17217	20398	26936	38444	38690	49698	49700	68299	70940	8523	8662	9998	86558	437	51487	51195	51188	51140	51541	51186	51052	60722	98609
	Left 86763	87040	85824	86844	110731	110763	110923	84896	101610	102591	13919	14463	16216	17210	20385	22177	38443	38677	49697	49699	19199	70934	8497	8661	8665	86557	436	51485	51193	51186	51138	51539	51184	51050	60720	60984
Marker	Name 474374	466971	466972	466973	467792	467793	467794	469174	469658	469659	469660	469661	469662	469663	469664	469665	469666	469667	469668	469669	469670	469671	469672	469673	469674	471147	470500	471938	471939	471940	471941	471942	471943	471944	473070	473071
BAC	Name T14C9	F15F15	F15F15	F15F15	F15F15	F15F15	F15F15	F15F15	F15F15	F15F15	F15F15	F15F15	F15F15	F15F15	F15F15	F15F15	F15F15	MPK17	F16F17																	
BAC	Chromosome Length	157000	157000	157000	157000	157000	157000	157000	157000	157000	157000	157000	157000	157000	157000	157000	157000	157000	157000	157000	157000	157000	157000	157000	157000	157000	16898	61510	61510	61510	61510	61510	61510	61510	61510	61510
	Chromosoi 5	ט י	5	5	5	5	S	5	5	5	S	5	5	\$	5	5	5	5	5	5	5	5	5	5	S	5	5	5	S	5	S	S	S	5	5	5
	Seq id	AC007627	AC007627	AC007627	AC007627	AC007627	AC007627	AC007627	AC007627	AC007627	AC007627	AC007627	AC007627	AC007627	AC007627	AC007627	AC007627	AP000418	AB028606																	
Seq	num 120	121	121	121	121	121	121	121	121	121	121	121	121	121	121	121	121	121	121	121	121	121	121	121	121	121	122	123	123	123	123	123	123	123	123	123

NP Base Columbia	Landsberg																					
		O/C	CI	T/A	T/A	T/G	A/T	G/T	G/A	G/A	A/C	A/T	T/A	G/A	T/A	G/A	T/A	A/C	A/G	A/G	A/T	T/A
Indel Size Columbia/	Landsberg																					
	Method	_	-1	1	1				_	-	-	1	-	T	1	1	1	-		_		-
	Type	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP	SNP
	Right	<i>LLL</i> 2011 100 100 100 100 100 100 100 100 10	61178	36128	36088	36046	36129	36069	26416	26321	26248	26465	33848	105	159	383	720	153	364	16841	14094	1817
	Left	60775	61176	36126	36086	36044	36127	36067	26414	26319	26246	26463	33846	103	157	381	718	151	362	16839	14092	1815
Marker	Name	473072	473073	471929	471930	471931	471932	471933	472117	472118	472119	472120	472434	472550	472551	472552	472553	472554	472555	473085	473089	473444
BAC	Name	F16F17	F16F17	F10E10																		
BAC	Chromosome Length	61510	61510	38089	38089	38089	38089	38089	38089	38089	38089	38089	38086	38089	38089	38089	38089	38088	38089	38089	38089	38089
	Chr	3	5	S	S	5	3	Ŋ	5	v	, v	v	v	·	v	, v	v	, v	v	٠,	S	5
	Seq id	AB028606	AB028606	AB028605	VB028605	AB028605																
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## Example 3

SNPs are identified by comparing Arabidopsis thaliana, Columbia and Arabidopsis thaliana, Landsberg erecta sequences. Each Arabidopsis thaliana, Columbia BAC sequence (extracted from GenBank and represented by a SEQ ID NO: 1 through SEQ ID NO: 124) is compared to a full set of Arabidopsis thaliana, Landsberg erecta contigs using WUBLAST (version 2.0) to locate areas of high identity that could contain a marker. Each identified contig is subsequently compared using WUBLAST to a full set of Arabidopsis thaliana, Columbia BACs (all of SEQ ID NO: 1 through SEQ ID NO: 124). To be selected as a marker candidate, an Arabidopsis thaliana, Landsberg erecta contig must have either one or two matches to an Arabidopsis thaliana, Columbia BAC. A single match suggests that that the sequence is unique. Two matches often result from overlapping BACs. The alignments are evaluated in a conservative manner. False negatives are preferable to false positives. To be included as a candidate polymorphic marker there must be: a minimum alignment of 200 bases between the sequence of an Arabidopsis thaliana, Landsberg erecta contig and the sequence of an Arabidopsis thaliana, Columbia BAC; the alignment must cover at least 75% of the length of the Arabidopsis thaliana, Landsberg erecta contig; a minimum of two reads of the Arabidopsis thaliana, Landsberg erecta region with the two read areas extending at least 25 bases on each side of the polymorphism position; agreement between all Arabidopsis thaliana, Landsberg erecta reads at the polymorphism position; minimum PHRAP consensus quality of 40 at the polymorphism position, with an average quality of 30 for the 25 bases on each side of the polymorphism position; and a maximum 1% polymorphism across the sequence. SNPs and INDELs of less than three nucleotide bases identified as described above are set forth in Table A.

A set of fifty polymorphisms was selected from among the polymorphisms in Table A.

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## Example 4

PCR primers can be designed for the flanking sequence of polymorphisms and can be used to either confirm or detect the polymorphisms. Such primers are designed with the program Primer3 (obtained from the MIT-Whitehead Genome Center) with a "perl-oracle" wrapper. The criteria applied to design a primer include:

Primer annealing temperature (minimum 57°C, optimum 60°C, maximum 63 °C)

Primer length (minimum 18 bp, optimum 20 bp, maximum 27 bp)

G+C content (minimum 20%, maximum 80%)

Minimum target margin of the primer relative to the polymorphism: 50 bp

10 Length of the amplified region

for SNPs: minimum 480 bp, optimum 500 bp, maximum 550 bp for INDELs: minimum 200 bp, optimum 400 bp, maximum 500 bp

PHRED quality score of the gene template (minimum of 0)

Target sequence on one contig

15 Maximum mismatch = 12.0 (weighted score from Primer3 program)

Pair Max Misprime = 24.0 (weighted score from Primer3 program)

Maximum N's = 0

Maximum poly-X = 5

The primary goal of the design process is the creation of groups of primer pairs with a common annealing temperature  $(T_m)$ .

After the *Arabidopsis thaliana* specific portion of the primers is selected, an additional common primer tail sequence can be added to the 5' ends. Forward primers for the detection of insertion/deletion polymorphisms have the additional common M13 bases on the 5' end: (5'-CAGCACGTTGTAAAACGAC-3'); reverse primers for the detection of insertion/deletion polymorphisms were designed without a tail. Forward primers for the detection of SNPs have the additional common M13 bases on the 5' end: (5'-TGTAAAACGACGGCCAGTT-3'); reverse primers for SNPs have the additional

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common M13 bases on the 5' end: (5'-CAGGAAACAGCTATGACC-3'). The primer tail sequences are added so that subsequent amplifications of any primer pair can be done with a specific kit designed to work with oligonucleotides having the primer tail. It is noted that primer pairs are not required to contain the tail sequence, the relevant portion for amplification and/or hybridization probes being the *Arabidopsis thaliana* specific sequences.

Using such primers for polymorphic marker flanking sequence, a person skilled in the art can amplify genetic regions from Arabidopsis thaliana, Columbia and Arabidopsis thaliana, Landsberg erecta genomic DNA, as well as from a mixture of Arabidopsis thaliana, Columbia and Arabidopsis thaliana, Landsberg erecta genomic DNA to represent a heterozygote. In the case of SNPs the amplified product is purified and sequenced to confirm the presence of a predicted SNP. For validation of INDELs, the amplified products are analyzed or sized on an agarose gel or an acrylamide gel to determine if the fragments amplified from Arabidopsis thaliana, Columbia and Arabidopsis thaliana, Landsberg erecta genomic DNA are polymorphic. An exemplary PCR amplification reaction procedure to detect an INDEL-type polymorphism in a mapping experiment is a follows: a reaction mixture containing 4 ng/µl DNA (2.6µl); Taq Gold Polymerase (5 units/μl) (0.1μl) (Perkin Elmer, Norwalk, Connecticut); 5 μm forward and reverse primer (0.2µl); 1µm Li-Cor M13 Forward/IRD 700 (0.5µl)(Lincoln, Nebraska); 50 mM MgCl<sub>2</sub> (0.3μl); 10 mM dNTPs (2.5 mM each of dCTP, dGTP, dATP and dTTP)(0.8µl); 10X Taq Gold Buffer (1.0µl); dH<sub>2</sub>O (4.5µl). Thermal amplification is carried out in an MJ Tetrad as follows: 94°C 10 minutes; 35 cycles (94°C 1 minute, 56°C 1 minute, 72°C 1 minute); 72°C 10 minutes; 4°C hold. PCR products are loaded on a 7% Long Ranger gel and run on Li-Cor's DNA Sequencer Long Redir 4200 or DNA Analyzer Gene Readir 4200 according to manufacturer's protocol. Data is analyzed using GeneImagIR software.

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An exemplary PCR amplification reaction to detect a SNP-type polymorphism in a mapping experiment is a follows: A reaction mixture containing 4ng/µl DNA (6.6µl); 5 units Platinum Gold Polymerase (5 units/µl)(0.1µl) (GibcoBRL, Rockville, Maryland (0.11µl); 5 µm forward and reverse primer with M13 tails (1.39µl); 50 mM MgCl<sub>2</sub> (0.66µl); 10 mM dNTPs (2.5 mM each of dCTP, dGTP, dATP and dTTP)(1.04µl); 10X Taq Platinum Buffer (2.43µl); dH<sub>2</sub>O (12.77µl). Thermal amplification is carried out in an MJ Tetrad as follows: 94°C 10 minutes; 35 cycles (94°C 1 minute, 56°C 1 minute, 72°C 1 minute); 72°C 10 minutes; 4°C hold. PCR products are purified using QIAGEN's QIAquick 96 PCR Purification Kit as per manufactures' protocol. Purified PCR products are run on agarose gels to confirm amplification, followed by sequencing to confirm the presences of a SNP.

## We claim:

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- 1. A method of isolating a region of genomic DNA associated with a phenotype of interest comprising:
- (A) identifying an *Arabidopsis* plant of a first ecotype with a phenotype of interest;
  - (B) crossing said *Arabidopsis* plant with an *Arabidopsis* plant of a second ecotype lacking said phenotype;
    - (C) propagating and self pollinating seeds from said cross;
    - (D) selecting progeny of self pollinated seeds with said phenotype;
- (E) screening progeny of self pollinated seeds with said phenotype with a collection of nucleic acid molecules, said collection of nucleic acid molecules capable of detecting a set of polymorphisms where the polymorphisms are distributed throughout the genome of said self pollinated seeds with said phenotype at an average density of more than one polymorphism per about 100kb, wherein at least one of the polymorphisms is selected from Table A;
- (F) calculating the linkage of each of said polymorphisms to said phenotype; and
- (G) isolating said region of genomic DNA associated with said phenotype based on its linkage to one or more of said nucleic acid molecules.
- 2. The method of isolating a region of genomic DNA associated with a phenotype of interest according to claim 1, wherein said region of genomic DNA associated with said phenotype is located between about 5 and about 10 cM of one or more of said polymorphisms.
- 3. The method of isolating a region of genomic DNA associated with a phenotype of interest according to claim 1, wherein said region of genomic DNA associated with said

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phenotype is located between about 0 and about 5 cM of one or more of said polymorphisms.

- 4. A method of identifying a region of genomic DNA associated with a phenotypic trait of interest comprising:
  - (A) screening a mapping population of *Arabidopsis* plants to determine the linkage of said phenotypic trait with a collection of nucleic acid molecules, wherein said nucleic acid molecules are capable of detecting a set of polymorphisms, where the polymorphisms are distributed throughout the genome of said mapping population of *Arabidopsis* plants at an average density of more than one polymorphism per about 100kb, wherein at least one of the polymorphisms is selected from Table A;
  - (B) calculating the linkage of each of said polymorphisms to said phenotypic trait; and
- (C) identifying said genomic DNA region associated said phenotypic trait based on its linkage to one or more of said nucleic acid molecules.
  - 5. The method of identifying a region of genomic DNA associated with a phenotypic trait of interest according to claim 4, further comprising isolating said identified region.
- 20 6. The method of identifying a region of genomic DNA associated with a phenotypic trait of interest according to claim 4, wherein said collection of nucleic acid molecules is capable of detecting a set of greater than 25 polymorphisms selected from Table A.
- 7. The method of identifying a region of genomic DNA associated with a phenotypic trait of interest according to claim 6, wherein said collection of nucleic acid molecules is capable of detecting a set of greater than 50 polymorphisms selected from Table A.

- 8. The method of identifying a region of genomic DNA associated with a phenotypic trait of interest according to claim 7, wherein said collection of nucleic acid molecules is capable of detecting a set of greater than 75 polymorphism selected from Table A.
- 5 9. The method of identifying a region of genomic DNA associated with a phenotypic trait of interest according to claim 8, wherein said collection of nucleic acid molecules is capable of detecting a set of greater than 100 polymorphisms selected from Table A.
- 10. A method of identifying a nucleic acid molecule associated with a phenotypic trait10 of interest comprising:
  - (A) screening a mapping population of *Arabidopsis* plants to determine the linkage of said phenotypic trait with a collection of polymorphisms, wherein said polymorphisms are distributed throughout the genome of said mapping population of *Arabidopsis* plants at an average density of more than one polymorphism per about 100kb, wherein at least one of the polymorphisms is selected from Table A;
  - (B) calculating the linkage of each of said polymorphism to said phenotypic trait; and
  - (C) isolating said nucleic acid molecule associated with said phenotypic trait based on its linkage to one or more of said polymorphisms.
  - 11. The method of identifying a nucleic acid molecule associated with a phenotypic trait of interest according to claim 10, wherein said collection of polymorphisms comprises at least 25 polymorphisms selected from Table A.
- 25 12. The method of identifying a nucleic acid molecule associated with a phenotypic trait of interest according to claim 11, wherein said collection of polymorphisms comprises at least 50 polymorphisms selected from Table A.

13. The method of identifying a nucleic acid molecule associated with a phenotypic trait of interest according to claim 12, wherein said collection of polymorphisms comprises at least 75 polymorphisms selected from Table A.

- 14. The method of identifying a nucleic acid molecule associated with a phenotypic trait of interest according to claim 13, wherein said collection of polymorphisms comprises at least 100 polymorphisms selected from Table A.
- 15. The method of identifying a nucleic acid molecule associated with a phenotypic trait of interest according to claim 10, wherein said nucleic acid molecule associated with said phenotypic trait is located between about 5 and about 10 cM of one or more of said polymorphisms.
- 15 16. The method of identifying a nucleic acid molecule associated with a phenotypic trait of interest according to claim 15, wherein said nucleic acid molecule associated with said phenotypic trait is located between about 0 and about 5 cM of one or more of said polymorphisms.
- 20 17. A method of isolating a nucleic acid molecule associated with a phenotypic trait comprising:
  - (A) screening a mapping population of *Arabidopsis* plants to determine the linkage of said phenotypic trait with a collection of polymorphisms, wherein said at least one polymorphism is selected from Table A; and
- 25 (B) isolating said nucleic acid molecule associated with said phenotypic trait based on its linkage to one or more of said polymorphisms.

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- 18. A collection of non-identical nucleic acid molecules capable of detecting polymorphisms present in an *Arabidopsis* mapping population, wherein said collection of non-identical nucleic acid molecules is capable of detecting at least 25 polymorphisms selected from the group consisting of Table A.
- 19. The collection of non-identical nucleic acid molecules capable of detecting polymorphisms present in an *Arabidopsis* mapping population according to claim 18, wherein said collection non-identical nucleic acid molecules is capable of detecting at least 25 polymorphisms which are single nucleotide polymorphisms.
- 20. The collection of non-identical nucleic acid molecules capable of detecting polymorphisms present in an *Arabidopsis* mapping population according to claim 18, wherein said collection non-identical nucleic acid molecules is capable of detecting at least 25 polymorphisms are insertion or deletion polymorphisms.
- 21. The collection of non-identical nucleic acid molecules capable of detecting polymorphisms present in an *Arabidopsis* mapping population according to claim 518, wherein said collection of non-identical nucleic acid molecules is capable of detecting at least 50 polymorphisms selected from the group consisting of Table A.
- 22. The collection of non-identical nucleic acid molecules capable of detecting polymorphisms present in an *Arabidopsis* mapping population according to claim 21, wherein said collection of non-identical nucleic acid molecules is capable of detecting at least 100 polymorphisms selected from the group consisting of Table A.

23. The collection of non-identical nucleic acid molecules capable of detecting polymorphisms present in an *Arabidopsis* mapping population according to claim 18, wherein said collection of non-identical nucleic acid molecules is deposited on a substrate.

- 24. Computer readable medium having recorded thereon at least 100 of the polymorphisms set forth in Table A.
- 25. A method of introgressing a trait of interest into a plant comprising using a nucleic acid marker for marker assisted selection of said plant, said nucleic acid marker capable of detecting a polymorphism selected from Table A, and introgressing said trait into said plant.
- 26. A method for identifying transposons in the DNA of an organism comprising
   15 identifying INDELs in said DNA and comparing the sequence of said INDELs to the sequence of one or more known transposons.

## **Abstract**

The present invention is in the field of plant genetics. More specifically, the invention relates to nucleic acid markers associated with *Arabidopsis thaliana* ecotypes.

5 The invention also relates to methods for detecting polymorphisms. The invention further relates to methods of using nucleic acid markers, for example, for genome mapping, gene identification, gene isolation and gene analysis.